

**PCT**WORLD INTELLECTUAL PROPERTY ORGANIZATION  
International Bureau

## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

|  |           |   |
|--|-----------|---|
| <b>(51) International Patent Classification 5 :</b><br><b>A61K 31/70, 47/48</b>  | <b>A2</b> | <b>(11) International Publication Number:</b> <b>WO 90/10448</b><br><b>(43) International Publication Date:</b> 20 September 1990 (20.09.90)  |
| <b>(21) International Application Number:</b> PCT/US90/01002<br><b>(22) International Filing Date:</b> 23 February 1990 (23.02.90)<br><b>(30) Priority data:</b><br>320,202 7 March 1989 (07.03.89) US<br><b>(71) Applicant:</b> GENENTECH, INC. [US/US]; 460 Point San Bruno Boulevard, South San Francisco, CA 94080 (US).<br><b>(72) Inventors:</b> BISCHOFBERGER, Norbert, W. ; 1116 Cedar Street, San Carlos, CA 94070 (US). SHEA, Regan, G. ; 365 Laurie Meadows Drive No. 219, San Mateo, CA 94403 (US).<br><b>(74) Agents:</b> WINTER, Daryl, B. et al.; Genentech, Inc., 460 Point San Bruno Boulevard, South San Francisco, CA 94080 (US). |           | <b>(81) Designated States:</b> AT (European patent), BE (European patent), CA, CH (European patent), DE (European patent), DK (European patent), ES (European patent), FR (European patent), GB (European patent), IT (European patent), JP, LU (European patent), NL (European patent), SE (European patent).<br><br><b>Published</b><br><i>Without international search report and to be republished upon receipt of that report.</i> |
| <b>(54) Title:</b> COVALENT CONJUGATES OF LIPID AND OLIGONUCLEOTIDE<br><br><b>(57) Abstract</b><br><br>A conjugate is provided wherein a lipid is covalently coupled to an oligonucleotide having a nucleotide sequence that is either of interest or substantially complementary to a nucleotide sequence of interest. The conjugate is useful for nucleic acid purification, diagnostic methods, and therapeutic applications. The therapeutic aspect includes conjugates wherein the oligonucleotide contains an antisense nucleotide sequence to a sequence encoding a polypeptide responsible for a pathogenic disorder.                        |           |   |

**BEST AVAILABLE COPY**

*FOR THE PURPOSES OF INFORMATION ONLY*

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

|    |                              |    |  |    |                          |
|----|------------------------------|----|--|----|--------------------------|
| AT | Austria                      | ES | Spain                                    | MG | Madagascar               |
| AU | Australia                    | FI | Finland                                  | ML | Mali                     |
| BB | Barbados                     | FR | France                                   | MR | Mauritania               |
| BE | Belgium                      | GA | Gabon                                    | MW | Malawi                   |
| BF | Burkina Faso                 | GB | United Kingdom                           | NL | Netherlands              |
| BG | Bulgaria                     | HU | Hungary                                  | NO | Norway                   |
| BJ | Benin                        | IT | Italy                                    | RO | Romania                  |
| BR | Brazil                       | JP | Japan                                    | SD | Sudan                    |
| CA | Canada                       | KP | Democratic People's Republic<br>of Korea | SE | Sweden                   |
| CF | Central African Republic     | KR | Republic of Korea                        | SN | Senegal                  |
| CG | Congo                        | LI | Liechtenstein                            | SU | Soviet Union             |
| CH | Switzerland                  | LK | Sri Lanka                                | TD | Chad                     |
| CM | Cameroon                     | LU | Luxembourg                               | TG | Togo                     |
| DE | Germany, Federal Republic of | MC | Monaco                                   | US | United States of America |
| DK | Denmark                      |    |  |    |                          |

-1-

COVALENT CONJUGATES OF  
LIPID AND OLIGONUCLEOTIDE

5 This invention relates to covalent conjugates of lipids and oligonucleotides and their use in the pharmaceutical, purification, and diagnostic arts.

Use of antisense RNA or DNA has proven useful for manipulating eukaryotic gene expression. This technique is based on blocking the informational flow from DNA to protein via RNA by introducing a complementary sequence to a portion of the target mRNA in the cell. When the oligomer is absorbed by the cells, it can either function in the cytoplasm by blocking translation of the target messenger or, upon penetrating the nucleus, interfere with nuclear processing or transcription by binding to DNA. In the case of viral infection, antisense oligomers may also function by blocking viral replication, depending on the target site of the oligomer on the viral genome. Complementary nucleic acids that inactivate gene expression have been designated antisense nucleic acids.

15 Efficient transport of nucleic acids across cell membranes is of great importance for antisense nucleic acid technology, as well as in simple transfection experiments. For recent reviews of antisense oligonucleotides, see van der Krol *et al.*, BioTechniques, 6: 958 (1988) and Zon, Pharmaceutical Res., 5: 539 (1988).

Uptake of nucleotides by cells and liposomes is normally an inefficient process due to the high charge density of the nucleotide. A number of schemes have been used to improve incorporation efficiency, namely, calcium phosphate-mediated gene transfer (Chen, C.A. and Okayama, H., BioTechniques, 6:632 (1988)), use of retroviral vectors (Eglitis, M.A. *et al.*, BioTechniques, 6: 608 (1988)), microinjection (DePamphilis, M.L. *et al.*, BioTechniques, 6: 662 (1988)), use of adenovirus vectors (Berkner, K.L., BioTechniques, 6: 616 (1988)), electroporation (Andreason, G.L. and Evans, G.A., BioTechniques, 6: 650 (1988)), rapid acceleration of DNA-coated, gold particles (Christou P. *et al.*, Plant Physiol., 87: 671-674 (1988)), and liposome-mediated gene transfer (Mannino, R.J. and Gould-Fogerite, S., BioTechniques, 6: 682 (1988)). All of these methods, however, suffer from one or more problems relating to cellular toxicity, introduction of potentially pathogenic viral factors, poor reproducibility, inconvenience, or inefficiency of DNA delivery.

More recently, DNA covalently attached to cell receptor ligand proteins, such as that for epidermal growth factor, was studied as a means to effect cell-specific DNA delivery. EP 273,085 published July 6, 1988. Additionally, DNAs complexed with sugar-lysine conjugates (Wu, G.Y. and Wu, C.H., J. Biol. Chem., 263: 14621 (1988)) and with a positively charged lipid, i.e., N-[1-(2,3-dioleoyloxy)propyl]-N,N,N-trimethylammonium chloride, entrapped in a liposome (Felgner, P.L. *et al.*, Proc. Natl. Acad. Sci. USA, 84: 7413 (1987)), were shown to enhance transfection efficiency.

Protein-DNA conjugates are found in nature and are important in viral replication. Vartapetian and Bogdanov, A.A. Prog. Nucl. Acid Res. Mol. Biol., 34: 210 (1987). Non-natural hybrids have been prepared in which DNA is linked to targeting, cleaving, or reporter groups, including peptides, biotin, fluorescent dyes, and EDTA-Fe. Zuckermann

-2-

et al., J. Am. Chem. Soc., **110**: 6592 (1988); Haralambidis et al., Tetrahedron Lett., **28**: 5199 (1987); Connolly, Nucl. Acids Res., **15**: 3131 (1987); Agrawal et al., Nucl. Acids Res., **14**: 6227 (1986); Smith et al., Nucl. Acids Res., **13**: 2399 (1985); Moser and Dervan, Science, **238**: 645 (1987).

5        Phosphatidyl nucleosides are biosynthetic intermediates in the lipid metabolism and a number of derivatives thereof showed interesting biological activities. Shuto, S. *et al.*, Chem. Pharm. Bull., **36**: 209 (1988) (5'-(3-*sn*-phosphatidyl)nucleosides for antileukemic activity); U.S. Pat. No. 4,797,479 issued Jan. 10, 1989 to Shuto et al. (phospholipid-nucleoside complex prepared by reacting L-glycerophospholipid with nucleoside in the  
10       presence of phospholipase D for treatment of tumors); Matsushita, T. *et al.*, Cancer Res., **41**: 2707 (1981) (nucleoside 5'-diphosphate-L-1,2-dipalmitin derivatives of 1- $\beta$ -D-arabinofuranosylcytosine, 9- $\beta$ -D-arabinofuranosyladenine, and tubercidin for enhanced catabolic stability); Turcotte, J.G., *et al.*, Biochem. Biophys. Acta, **619**: 604 (1980) and **619**:  
15       619 (1980) (analogs of cytidine diphosphate diacylglycerol containing the 1- $\beta$ -D-arabinofuranosyl moiety for anticancer activity). A cholesterol-nucleoside conjugate was recently reported for application in liposomes. Hashida *et al.*, Chem. Pharm. Bull., **36**: 3186 (1988).

      Combinations of ether lipid analogs and DNA-interactive agents, i.e., adriamycin, 4-hydroperoxycyclophosphamide, and cisplatin, were found to enhance anti-tumor activity  
20       in an additive fashion. Nosedá *et al.*, Cancer Res., **48**: 1788-1791 (1988).

      Transfection methods applicable for *in vivo* therapeutic purposes, e.g., in which foreign sequences of active nucleic acids can be applied to a living organism to cure some deficiencies of the cell metabolism, have been developed. For example, Cheng *et al.*, Nucl. Acids Res., **11**: 654-669 (1983) reports that the construct of chloramphenicol-acetyltransferase gene bound to  $\alpha$  2-macroglobulin was internalized in 3T3-4 cells.  
25       However, no evidence was provided that the internalized DNA was capable of performing a function in the host cells. EP 273,085, published July 6, 1988, discloses linking a nucleic acid to a cell homing or targeting factor that promotes the penetration of the foreign nucleotides through the cell membrane, but has little specificity regarding cell recognition.  
30       Such targeting factors include low density lipoproteins, growth factors, viral antigens, and the B chain of toxins.

      The use of hydrophobic 5'-protecting groups for the HPLC purification of chemically synthesized oligodeoxynucleotides by solid support methods was recently investigated. Seliger and Schmidt, J. Chromatography **397**: 141-151 (1987); Schmidt *et al.*, Nucleosides  
35       and Nucleotides, **7**: 795-799 (1988).

      It is an object of the present invention to provide a vehicle for introducing DNA across cell membranes that is highly efficient.

      It is another object to provide a transfection technique for internalization of active oligo- and polynucleotides, e.g., antisense DNA that can hybridize to mRNA in the cell and  
40       thus inhibit cellular or viral functions.

-3-

It is yet another object to provide an agent that is more stable than DNA-lipid complexes and is a substrate for cellular lipases so that the molecule will be cleaved when it comes into contact with membrane-bound intracellular cytoplasmic enzymes to free the drug for use by the cell.

5 It is a further object to provide a liposome encapsulating such an agent.

It is still a further object to provide an assay for nucleic acids and antibodies incorporating the agent.

These and other objects will be apparent to one of ordinary skill in the art.

10 These objects are achieved by the provision of a covalent conjugate of a lipid and an oligonucleotide and pharmaceutically acceptable salts thereof. Preferably, the oligonucleotide has a nucleotide sequence sufficiently complementary to a pathogenic nucleic acid or an oncogene to hybridize thereto.

In another aspect, the invention provides a conjugate that is labeled and one that is immobilized on a solid support.

15 In further embodiments, the invention provides a method for introducing the above conjugate into a host cell comprising transfecting the host cell with the conjugate.

In a still further aspect, the invention supplies a composition comprising a pharmaceutically acceptable carrier and the conjugate, with the carrier preferably being a liposome.

20 In another aspect, the invention provides a method comprising administering to a plant, animal, or human suffering from a pathogenic condition an effective amount of the above composition.

In still another aspect, the invention provides a method for the assay of a nucleic acid having a predetermined nucleotide sequence in a sample comprising:

25 (a) providing the conjugate as a labeled conjugate that has a nucleotide sequence capable of hybridizing to the predetermined sequence;

(b) immobilizing the labeled conjugate on a support;

(c) contacting the sample with the immobilized conjugate under conditions that would cause hybridization of the nucleic acid with the oligonucleotide portion of the conjugate if the nucleic acid is present in the sample; and

30 (d) detecting the presence of labeled oligomers.

In an additional aspect, the invention provides a method for separating an oligonucleotide from a mixture, which method comprises:

(a) providing the conjugate having a nucleotide sequence capable of hybridizing to a sequence contained within the oligonucleotide to be separated;

35 (b) immobilizing the conjugate on a support;

(c) contacting the mixture with the immobilized conjugate, under conditions causing hybridization of the oligonucleotide of the mixture with the oligonucleotide portion of the conjugate; and

40 (d) separating the hybridized oligonucleotides.

-4-

In yet another aspect, the invention provides a method for separating an oligonucleotide from a mixture, which method comprises:

(a) providing the conjugate having a nucleotide sequence capable of hybridizing to a sequence contained within the oligonucleotide to be separated in a hydrophobic phase;

5 (b) providing the mixture in a hydrophilic phase;

(c) contacting the phase containing the mixture with the phase containing the conjugate, under conditions causing hybridization of the oligonucleotide of the mixture with the oligonucleotide portion of the conjugate; and

10 (d) extracting from the phase containing the mixture the hybridized oligonucleotides and transporting them to a separate hydrophilic phase.

In another aspect, the invention provides a method for detecting lupus erythematosus in a human comprising contacting a serum sample from the human with a liposome comprising the conjugate and a label, said contacting being such that the antibodies bind to the oligonucleotide on the liposome, so as to alter the stability of the liposome; and  
15 measuring for the presence or absence of the label.

The invention herein revolves around covalent conjugates of lipids and oligonucleotides that contain nucleotide sequences of interest or are sufficiently complementary to hybridize to sequences of interest. The lipid acts to internalize the nucleic acid sequence chemically coupled thereto into host cells, typically by endocytosis,  
20 and is bound to the oligonucleotide through a coupling sufficiently labile that it does not repress subsequent biochemical functioning of the oligonucleotide in the cell after internalization. In fact, preferably the conjugate is a substrate for a cellular enzyme that cleaves the conjugate to release the oligonucleotide. The lipid is suitably covalently bonded to any moiety on the oligonucleotide, such as an amino group on the base, the hydroxyl group of the phosphate, or a hydroxyl group at the 5' or 3' terminus of the oligonucleotide.  
25 Preferably, however, it is bonded to the 5' hydroxyl group thereof.

For purposes herein, the term "lipid" refers to fats and fat-derived materials that are insoluble in water but soluble in biologically occurring hydrophobic solvents, related to fatty acid esters, fatty alcohols, sterols, or waxes, and utilizable by the animal organism.  
30 Examples of such lipids include fatty acids and esters thereof, glycerides, e.g., triglycerides, glyceryl ethers, phospholipids, sphingolipids, fatty alcohols, waxes, terpenes, and steroids. Lipids include those that are derived naturally as well as those that are synthetically prepared.

Of particular interest herein are conjugates containing a cleavage site within the lipid  
35 or representing the covalent bond between the lipid and oligonucleotide that is specifically recognized by an enzyme endogenous to a host cell to which the conjugate is targeted. Thus, cleavage at the site will break the bond or hydrolyze the lipid so as to render the conjugate more water soluble. Preferably the cleavage site is susceptible to enzyme hydrolysis. Also preferred is that the enzyme is located at a cellular or nuclear membrane  
40 or is in the cytoplasm of a cell.

-5-

The most preferred lipids herein are phospholipids, where the enzyme that recognizes the cleavage site is a lipase, preferably a phospholipase. Also, preferably, the cleavage site is the bond between the 5'-hydroxyl oxygen atom of the oligonucleotide and the phosphorus atom (in the case of phospholipase D) or the bond between the oxygen atom on the glycerol side and the phosphorus atom (in the case of phospholipase C). Phospholipase C and D are ubiquitous membrane-bound cytoplasmic enzymes that catalyze the hydrolysis of certain phosphoester bonds, thereby cleaving the conjugate and releasing the oligonucleotide into the cytoplasm. For a review of phospholipases, see Dennis, E.A. in "The Enzymes," Vol. XVI (New York: Academic Press, Inc., 1983), p. 307-353). It is preferred to maximize cleavage by phospholipase C or D rather than other lipases that cleave fatty acids so as to leave lipid residue on the oligonucleotide.

Suitable phospholipids herein include, e.g., phosphoglycerides, plasmalogens, and other phosphatidic acids, sphingomyelin, and 3'-O-aminoacyl phosphatidyl glycerol.

The "oligonucleotide" portion of the conjugate refers to a molecule comprised of two or more deoxyribonucleotides or ribonucleotides that has a nucleotide sequence that is either of interest or sufficiently complementary to hybridize to a nucleotide sequence of interest. The oligonucleotide is typically one that is capable of performing a biochemical function in receptor host cells and/or altering the operation of the cell machinery. Its exact size will depend on many factors, which in turn depend on the ultimate function or use of the oligonucleotide. For example, if the oligonucleotide is to be used as an antisense oligomer, to ensure specificity thereof, the sequence must be unique in the mRNA population, and complementary to the target RNA.

It has been calculated that a chain length of 14 nucleotides may be sufficient to define the sequence specificity of oligonucleotides. See Ts'O, *et al.*, "Biological Approaches to the Controlled Delivery of Drugs" Vol. 507, Ann. N.Y. Acad. Sciences, 1987. Increasing the length of the oligonucleotide increases stability of the duplex formed and thus its inhibitory effect. On the other hand, very long oligomers can interact with two or more mRNAs through base pairing involving only 5-10 contiguous bases within the sequence of the oligonucleotide, thus lowering the specificity of the oligonucleotide. For general use, the oligonucleotide contains at least five bases, more preferably, from about five to about thirty bases, and most preferably from about fourteen to about twenty-five bases.

The oligonucleotides herein are selected either to contain, or to be sufficiently complementary to hybridize to, the nucleotide sequence of interest. Therefore, the oligonucleotide sequence need not reflect the exact sequence of the nucleotide sequence of interest. For example, a non-complementary nucleotide fragment may be attached to the 5' end of the oligonucleotide, with the remainder of the sequence being complementary to the nucleotide strand of interest. Alternatively, non-complementary bases or longer sequences can be interspersed into the oligonucleotide, provided that its sequence be capable of hybridizing to the target nucleic acid.

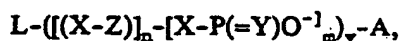
For purposes herein, the term "oligonucleotide" also includes oligonucleotides having modifications of the sugar-phosphodiester backbone to reduce their sensitivity to cellular

-6-

nucleases and increase absorption by the cell as needed. One example is methylphosphonates wherein one of the oxygen atoms is substituted by a methyl group to result in four different substituents on the phosphorus atom, as disclosed by van der Krol *et al.*, *supra* in Table 1. Oligomer methylphosphonates directed against HSV-1 immediate early mRNA-4 and -5 were found to prevent expression of herpetic lesions when applied in the form of cream to the HSV-infected ear of a mouse and were not toxic to mice when injected intravenously in concentrations up to 40 mg/kg body weight. Miller *et al.*, Anti-Cancer Drug Design, 2: 117-128 (1987); Ts'O *et al.*, *supra*. Additionally, phosphorothioates have been found to be potent inhibitors of viral proliferation, particularly HIV. Matsukura *et al.*, Proc. Natl. Acad. Sci., U.S.A., 84: 7706-7710 (1987); Agrawal *et al.*, Proc. Natl. Acad. Sci. U.S.A., 85: 7079-7083 (1988).

The definition of "oligonucleotides" also includes those with a covalently linked reagent besides the lipid that increases affinity for the complementary sequence. For example, coupling of poly-(L-lysine) to the 3' end of an antisense VSV oligonucleotide may be employed to enhance its antiviral activity. Further, a phosphorothioate oligomer may be coupled to a poly-(L-lysine) to lower the effective dose. In addition, a phosphorothioate oligonucleotide is useful herein because it is stable to cleavage by nucleases, is very soluble in water, and hybridizes more efficiently with a complementary DNA sequence than the corresponding methylphosphonate analogs. See, e.g., Marcus-Sekura *et al.*, Nucl. Acids Res., 15: 5749-5763 (1987). An intercalating agent such as acridine may be added to the 3' end of the oligonucleotide to enhance its affinity for the target. Other methods include attaching to the antisense oligomer a reactive agent to modify the target nucleic acid irreversibly, including alkylating reagents, metal complexes such as EDTA-Fe(II), o-phenanthroline-Cu(I), or porphyrin-Fe(II). Such compounds generate hydroxyl radicals in the presence of molecular oxygen and a reducing agent and cleave the complementary strand following attack on the target nucleic acid backbone. In addition, a photocrosslinking agent can be attached to the oligomer, such as a psoralen derivative, azidophenacyl, and proflavine.

In a preferred embodiment, the conjugate has the formula:



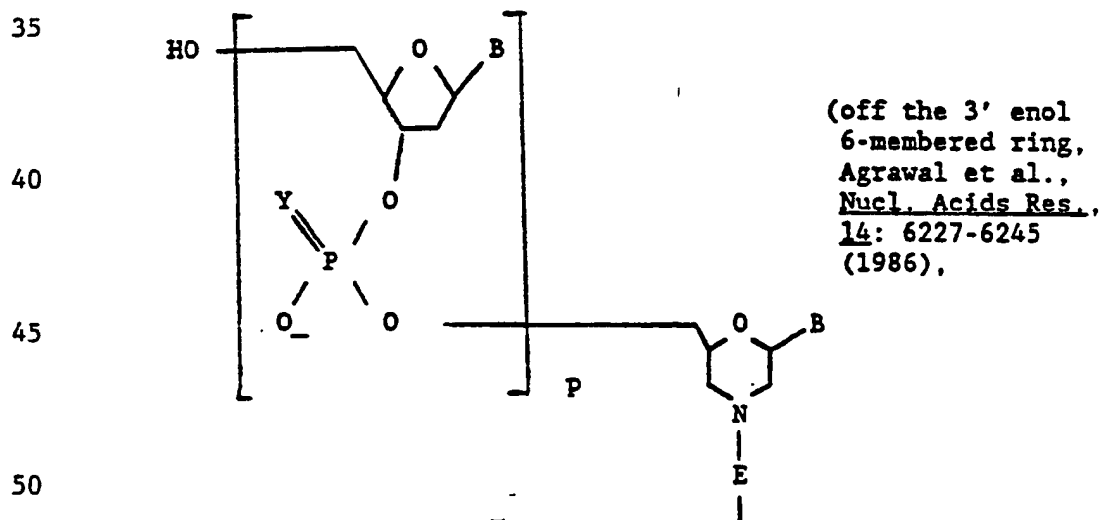
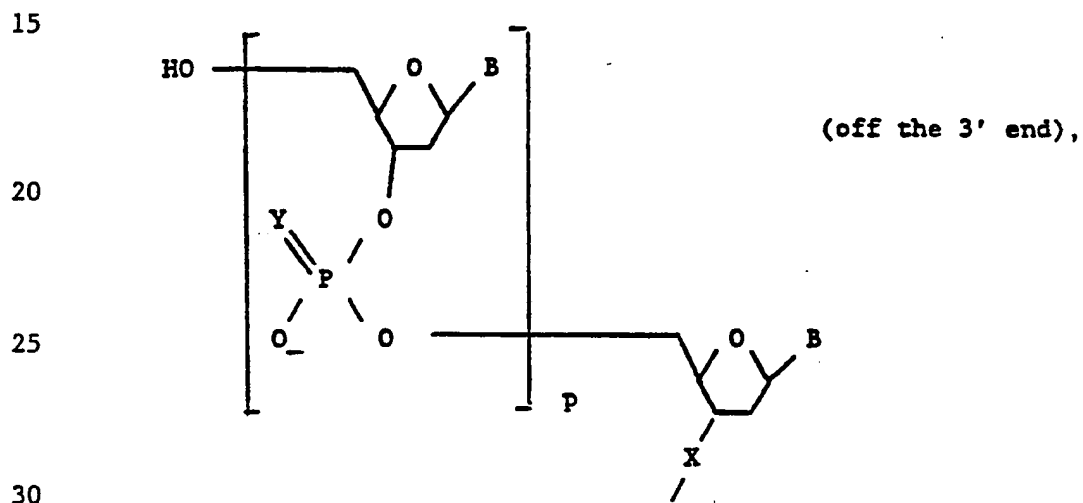
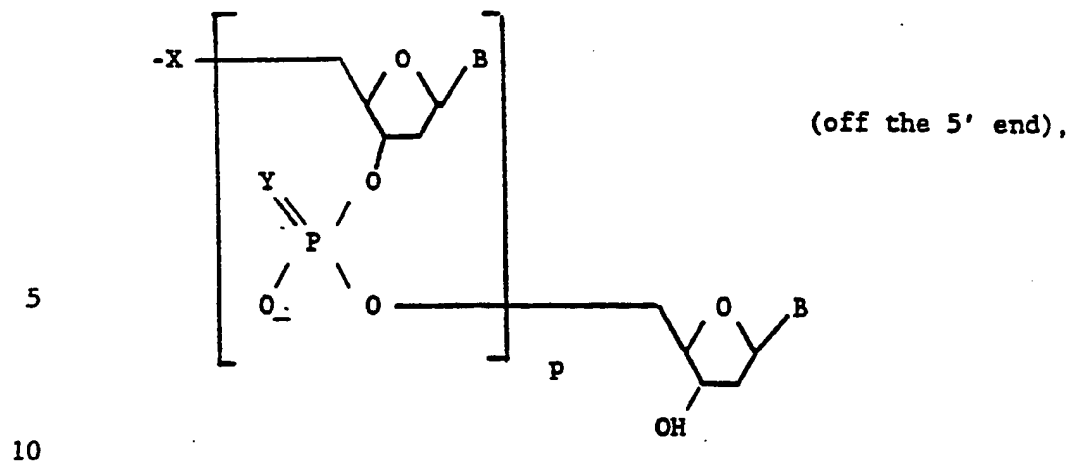
wherein L is a steroid moiety,  $R_1$ , or  $R_2-X-CH(R_4)-CH(-X-R_3)-CH_2$ , where  $R_1$  is a  $C_1$ - $C_{30}$  alkyl,  $C_2$ - $C_{30}$  mono-, di-, or polyunsaturated alkyl,  $C_3$ - $C_8$  cycloalkyl or  $C_4$ - $C_8$  mono-, di-, or polyunsaturated cycloalkyl group,  $R_2$ ,  $R_3$ , and  $R_4$  are independently H,  $R_1$ , or  $\alpha$ -amino acyl, X is O, S, NH, C(=O), or C(=O)O, OC(=O), NHC(=O), or C(=O)NH, and Y is O or S;

Z is a  $C_2$ - $C_{10}$  saturated or mono-, di- or polyunsaturated alkylene moiety, n, m, and y are independently an integer from 0 to about 10; and

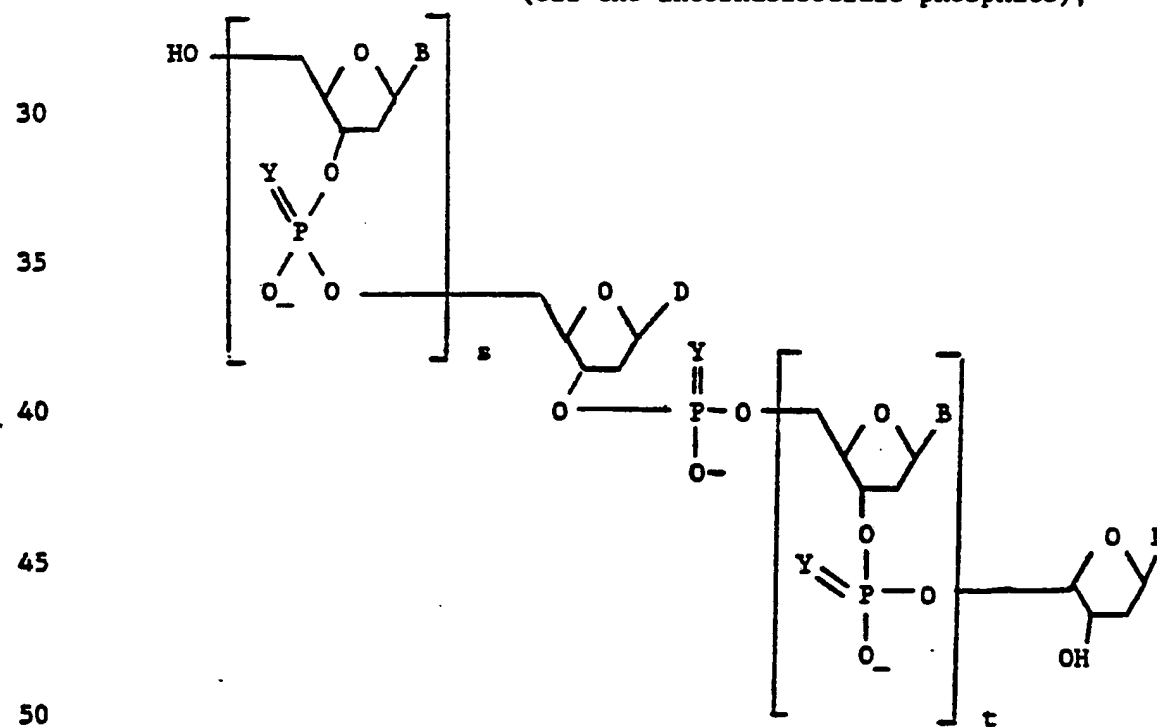
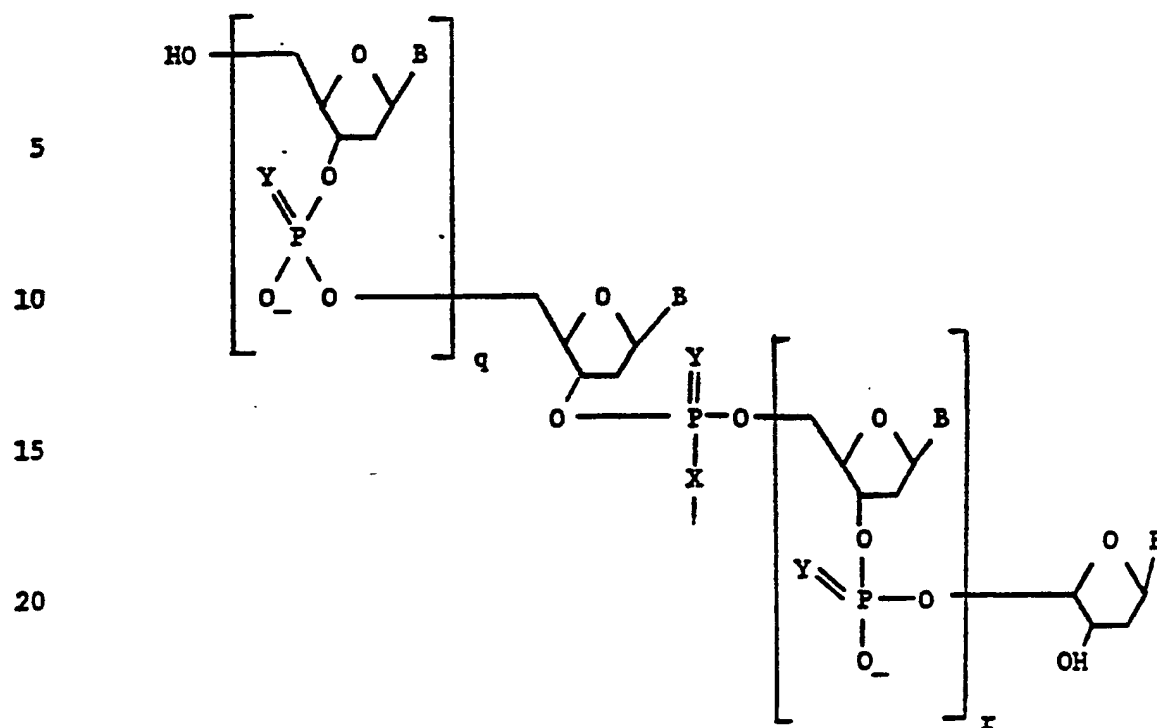
A is selected from the group consisting of



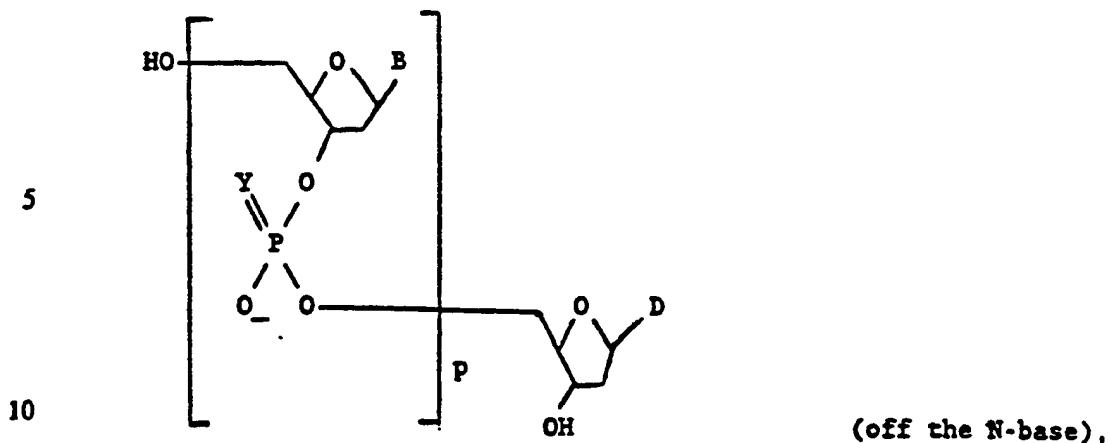
-7-



-8-

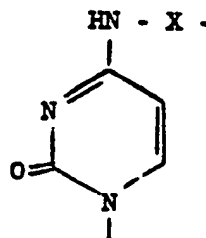
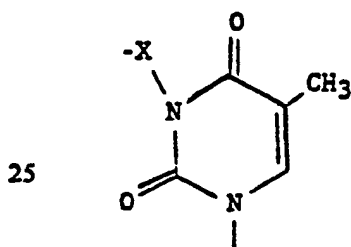


-9-

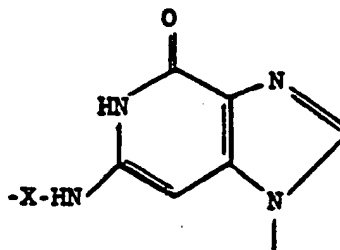
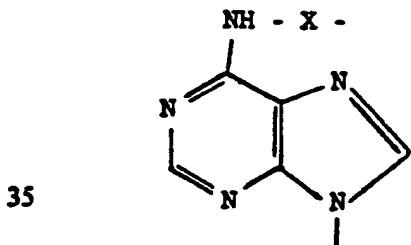


where B is a deprotected base, p is an integer from about 5 to 30, q and r are integers from about 1 to 28, provided that  $r + q$  is from about 4 to 29, s and t are integers from 0 to about 29, provided that  $s + t$  is from about 4 to 29, E is X or ZX, and D is selected from the group consisting of (where the bond from the ring nitrogens is attached to the sugar moiety of the oligonucleotide):

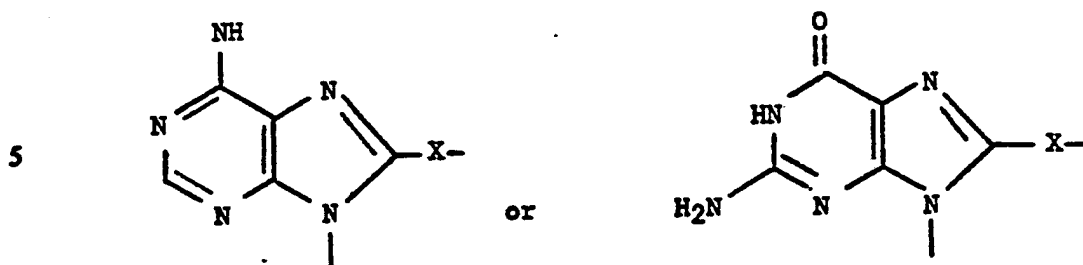
20



30



40



10 The terms "alkyl" and "cycloalkyl" as used herein refer to both straight-chain and branched alkyl groups and cycloalkyl groups with straight-chain or branched alkyl groups pending from the ring. "Mono-, di- and polyunsaturated" alkyl, cycloalkyl, or alkylene (divalent alkyl) moieties refer to hydrocarbons that contain one or more carbon-carbon double or triple bonds anywhere along their chain. Examples of compounds where L is an  
15 unsaturated cyclic or acyclic  $R_1$  include terpenes, such as phytol, geraniol, limonene, farnesol, and squalene, vitamins, such as vitamins A, D, and K, and  $\beta$ -carotene, and prostaglandins.

If L is a steroid moiety, preferably it is a sterol, and more preferably a sterol linked to the oligonucleotide through the hydroxyl group (either as an ether or ester) at its  $C_3$   
20 position. Examples of suitable steroids include cholesterol, lanosterol, a phytosterol, or a mycosterol such as, e.g., ergosterol.

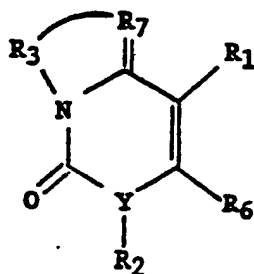
If L is  $R_2-X-CH(R_4)-CH(-X-R_3)-CH_2$ , then  $R_2$ ,  $R_3$ , and  $R_4$ , if defined as  $R_1$ , may be combined to form an additional carbon to carbon bond. Preferably,  $R_2$  and  $R_3$  are independently H or  $C_{10}$ - $C_{20}$  saturated or unsaturated alkyl groups,  $R_4$  is H or  $R_1$ , X is O,  
25 NH,  $NHC(=O)$ ,  $C(=O)$ ,  $OC(=O)$ , or  $C(=O)O$ , most preferably O, Y is O, Z is ethylene or carboxyethylene, n, m, and y are from 0 to 2, more preferably y is 1 or 2, m is 1 or 2, and n is 0 or 1, p is about 10 to 25, q and r are from about 1 to 23, provided that their sum is from about 9 to 24, and s and t are from 0 to about 24, provided that their sum is from about 9 to 24. Preferably A is the oligonucleotide connected through its 5' end, more  
30 preferably where X is O. If A is connected off the N-base, it is preferably off the N6 of an adenine residue or the N4 of a cytosine residue. If  $R_2$ ,  $R_3$ , and/or  $R_4$ , preferably  $R_2$ , is an  $\alpha$ -amino acyl moiety, preferably H is derived from any amino acid, more preferably a naturally occurring amino acid, and most preferably lysine.

If it is desired to provide oligonucleotides containing intercalating bases that are  
35 sterically confined so as to inactivate translationally the targeted strand, at least one of the bases A in the formula above is a substantially planar base. The oligonucleotides so derivatized hybridize to their complementary RNA or DNA strand, the modified nucleoside remaining unpaired but nonetheless intercalated between adjacent base pairs in the duplex in a precisely stereochemically defined manner. In light of the precise steric targeting made  
40 possible by this invention, the intercalating moiety is substituted with a reactive group capable of covalently modifying a predetermined site in the complementary domain. Such

-11-

reactive groups include crosslinking agents and phosphate bond cleaving agents. These reactive groups are sterically confined and less likely to interact with cellular components or nucleic acid at sites other than the target complementary sequence.

The phrase "substantially planar" in this context means that the steric bulk of the group lies substantially within an envelope approximating the steric gap bounded by the sugar backbone and flanking bases present in a complementary nucleic acid strand. In general, this envelope has dimensions of about 30-50 Angstroms in width and depth and about 3-7 Angstroms in thickness. An example includes a nucleoside having the structure:



wherein  $R_3$  is an aromatic polycycle, Y is C or N,  $R_7$  is N or  $=C(R_1)-$ ,  $R_1$  and  $R_6$  are H or a halogen, nitro, alkyl, hydroxyalkyl, or alkylether group wherein the alkyl group has 1 to 10 carbon atoms, and  $R_2$  is a ribose or deoxyribose sugar.

The nucleic acid sequences of interest include mRNA and DNA, and may be present in restriction enzyme digests or other fragments of nucleic acids, e.g., restriction fragment length polymorphisms (RFLPs). The nucleic acid or acids may be obtained from any source, for example, from plasmids such as pBR322, from cloned DNA or RNA, or from natural DNA or RNA from any source, including bacteria, yeast, viruses, and higher organisms such as plants or animals. For therapeutic purposes, the antisense oligonucleotides should be selected so as to have an efficient and specific interaction with the target mRNA, an efficient cellular uptake and compartmentalization of the oligonucleotide, and sufficient stability in the different cellular compartments.

For diagnostic use, the oligonucleotide may contain a sequence that encodes a diagnostically useful protein, e.g., pathogenic proteins, such as those responsible for viral infections, including AIDS, oncogenes, growth factors,  $\beta$ -globin, and the like. Further, other nucleic acids that encode no protein may be of interest, e.g., transcription or translation control domains or sequences useful in forensic medicine.

Most preferably, the conjugates herein have utility as agents for the antisense inhibition of translation of target nucleic acids. Such utilities have already been explored extensively with other antisense oligonucleotides (see van der Krol *et al.*, *supra*, and WO 83/01451 published April 28, 1983), and the oligonucleotides herein will be used in substantially the same fashion, using a rational, specific design based on the sequence and secondary structure information of the target RNA or DNA, keeping in mind that the

-12-

secondary or tertiary structures in the antisense RNA may influence the extent or rate of hybridization. For this use, the oligonucleotide has a nucleotide sequence capable of hybridizing to a pathogenic nucleic acid or an oncogene, more preferably viral or parasitic nucleic acid infecting a plant, such as a vegetable plant; animal, such as domestic, sports, and farm animals, e. g., dogs, cats, cows, and pigs; or human. For example, antisense oligonucleotides, either modified or unmodified, can be used to inhibit viral infections of Rous sarcoma virus, vesicular stomatitis virus, type A influenza virus, herpes simplex virus, human immunodeficiency virus, and plant viruses such as potato virus X coat protein gene and cucumber mosaic virus coat protein gene, to inhibit parasitic, e.g., malarial, infections, and to reduce expression of proto-oncogene *c-myc* (for leukemia treatment), simian sarcoma virus,  $\beta$ -globin, and  $\beta$ -tubulin genes. Examples of published viral sequences that have inhibitory effects include those against Rous sarcoma virus (Zamecnik and Stephenson, Proc. Natl. Acad. Sci. U.S.A., 75: 280-284 (1978); Stephenson and Zamecnik, Proc. Natl. Acad. Sci. U.S.A., 75: 285-288 (1978)), those against human T-cell lymphotropic virus type III (Zamecnik et al., Proc. Natl. Acad. Sci. U.S.A., 83: 4143-4146 (1986); WO 87/07300 published Dec. 3, 1987), and those against herpes simplex virus type 1 (Smith et al., Proc. Natl. Acad. Sci. U.S.A., 83: 2787-2791 (1986)).

Included within the definition of useful oligonucleotides for inactivating pathogenic nucleic acids are oligonucleoside alkyl- or arylphosphonate analogues complementary to the target pathogenic sequence and including a functional group that reacts with the target nucleic acid to render it inactive or nonfunctional. These derivatives are described in EP 266,099 published May 4, 1988.

In other embodiments, the oligonucleotide has a nucleotide sequence representing, or capable of hybridizing to, a cleavage site specifically recognized by an enzyme endogenous to a host cell to which the conjugate is targeted. For example, the oligonucleotide may encompass a sequence recognized by the RNase-H enzyme. This enzyme cleaves the RNA part of an RNA/DNA hybrid *in vivo*, resulting in subsequent degradation of the mRNA. Thus, the effect of the antisense oligomer can be catalytically enhanced by RNase-H activity.

Since the splice junctions of pre-mRNAs interact with the RNAs of small ribonucleoprotein particles that mediate the splicing process, these regions of the messenger are ideal targets for complementary oligomers. Thus, the invention herein includes antisense DNA or their analogs directed against splice-junction sites, especially when such antisense DNA has been found to be effective in inhibiting proliferation of HIV (Agrawal et al., Proc. Natl. Acad. Sci. U.S.A., 85: 7079-7083 (1988)), HSV (Smith et al., Proc. Natl. Acad. Sci. U.S.A., 83: 2787-2791 (1986)), or synthesis of SV40 large T antigen (Verspieren et al., Gene, 61: 307-315 (1987)). In addition, the invention is directed to complementary oligomers that are designed to interact with the cap site or the initiation codon region on the mRNA. Also, it has been shown that antisense oligomers blocking the ribosome attachment site are very effective for this purpose.

-13-

The oligonucleotides herein are suitably prepared using conventional methods such as, for example, the phosphotriester or phosphodiester methods described by Narang, S.A. *et al.*, Meth. Enzymol., 68: 90 (1979) and Brown, E.L. *et al.*, Meth. Enzymol., 68: 109 (1979), respectively, or automated embodiments thereof. In one such automated embodiment, phosphoramidites are used as starting materials and may be synthesized as described by Beaucage *et al.*, Tetrahedron Letters, 22: 1859-1862 (1981). One method for synthesizing oligonucleotides on a modified solid support is described in U.S. Pat. No. 4,458,066, issued July 3, 1984. It is also possible to use an oligonucleotide that has been isolated from a biological source, such as a restriction endonuclease digest.

The conjugates are prepared by any suitable technique. For example, H-phosphonate methodology may be used to synthesize phospholipid conjugates, as described by Lindh and Stawinski, J. Nucleic Acids Res., Symp. Ser. No. 18, 189 (1987). Synthesis of the same compounds is also accomplished using phosphoramidite chemistry as described in Beaucage and Caruthers, Tetrahedron Lett., 1859 (1981) and McBride and Caruthers, Tetrahedron Lett., 245 (1983). The main advantage of using the H-phosphonate methodology is the flexibility in the oxidation procedure. When the oxidation is carried out with carbon tetrachloride in the presence of amines, phosphoroamidates are available, whereas oxidation with sulfur leads to phosphorothioate analogs, and carbon tetrachloride in the presence of alcohols produces phosphate triesters. Froehler, Tetrahedron Lett., 27: 5575 (1986).

Chemical attachment of lipids to oligodeoxynucleotides by the standard solid support DNA synthesis protocol can be employed for lipids that are normally stable to the deblocking conditions, i.e., concentrated ammonia. Such lipids include ether lipids or amide lipids, including triglycerides and sterols that contain amide functionalities in their backbone. Fatty acid derived lipids that contain ester linkages would not normally be stable to such conditions. The latter lipids can be conjugated by using the method of Chu *et al.*, Nucl. Acids Res., 11: 6513-6529 (1983). This method requires that the lipid contain a free primary or secondary amino group. The lipid may already contain such an amino group in a side chain, as, e.g., in the case of phosphatidyl ethanolamine, phosphatidyl 3'-O-aminoacyl glycerol, and phosphatidyl serine, or it may be derivatized to contain such an amino group. Using the Chu *et al.* method, the aminolipid is coupled to an oligomer-5'-phosphate so as to produce lipid oligonucleotide conjugates. Also, phospholipids may be suitably bonded to oligonucleotides enzymatically using phospholipase D, as has been reported for the synthesis of nucleoside lipid conjugates. See Shuto *et al.*, *supra*.

One use for the conjugate herein is for the transfection of host cells so that DNA is stably incorporated therein. The transfection involves merely contacting the cells with the conjugate as by use of stirring at room temperature for at least about 15 hours, preferably about 20-30 hours.

Another use is for the treatment of a mammal with a pathogenic condition that is alleviated or cured by use of an oligonucleotide with a nucleotide sequence that blocks a nucleic acid responsible for the condition. Examples of such conditions include those caused by oncogenes and infections caused by viruses such as AIDS, herpes, hepatitis, etc.

-14-

The mammal to be treated may be any mammalian species such as domestic and farm animals, including primates, and sports or pet animals, as well as humans. Preferably, however, the preferred species being treated is human.

5       The conjugate is administered to the patient by any suitable technique, including parenteral, sublingual, topical, intrapulmonary, and intranasal administration. The specific route of administration will depend, e.g., on the type of therapy required. Examples of parenteral administration include intramuscular, subcutaneous, intravenous, intraarterial, and intraperitoneal administration.

10       The conjugates to be used in the therapy will be formulated and dosed in a fashion consistent with good medical practice taking into account the clinical condition of the individual patient, the cause of the condition in need of therapy, the site of delivery of the conjugate, the method of administration, the scheduling of administration, and other factors known to practitioners. The "effective amount" for purposes herein is thus determined by such considerations.

---



-15-

a nonionic surfactant such as Tween, Pluronic, or PEG, e.g., Tween 80, in an amount of 0.04-0.05% (w/v), to increase its solubility.

Optionally other ingredients may be added such as antioxidants, e.g., ascorbic acid; low molecular weight (less than about ten residues) polypeptides, e.g., polyarginine or tripeptides; proteins, such as serum albumin, gelatin, or immunoglobulins; hydrophilic polymers such as polyvinylpyrrolidone; amino acids, such as glycine, glutamic acid, aspartic acid, or arginine; monosaccharides, disaccharides, and other carbohydrates including cellulose or its derivatives, glucose, mannose, or dextrans; chelating agents such as EDTA; and sugar alcohols such as mannitol or sorbitol.

The conjugate to be used for therapeutic administration must be sterile. Sterility is readily accomplished by filtration through sterile filtration membranes (e.g., 0.2 micron membranes). The conjugate ordinarily will be stored in lyophilized form or as an aqueous solution or emulsion.

Any reference to the conjugates herein also includes the pharmaceutically acceptable salts of such compounds, and it will be understood that use of certain of the foregoing excipients, carriers, or stabilizers will result in the formation of such salts. Examples of pharmaceutically acceptable salts include those of alkaline earths (e.g., sodium or magnesium), ammonium or  $NX_4^+$  (wherein X is  $C_{1-4}$  alkyl). Other pharmaceutically acceptable salts include organic carboxylic acids such as acetic, lactic, tartaric, malic, isethionic, lactobionic, and succinic acids; organic sulfonic acids such as methanesulfonic, ethanesulfonic, benzenesulfonic, and p-toluenesulfonic acids; and inorganic acids such as hydrochloric, sulfuric, phosphoric, and sulfamic acids. Physiologically acceptable salts of a compound having a hydroxy group include the anion of said compound in combination with a suitable cation such as  $Na^+$ ,  $NH_4^+$ , and  $NX_4^+$  (wherein X is a  $C_{1-4}$  alkyl group).

Therapeutic conjugate compositions generally are placed into a container having a sterile access port, for example, an intravenous solution bag or vial having a stopper pierceable by a hypodermic injection needle.

The conjugates herein are also suitably administered by sustained release systems. Suitable examples of sustained release compositions include semi-permeable polymer matrices in the form of shaped articles, e.g., films, or microcapsules. Sustained release matrices include polylactides (U.S. Pat. No. 3,773,919, EP 58,481), copolymers of L-glutamic acid and gamma-ethyl-L-glutamate (U. Sidman et al., *Biopolymers*, **22**, 547-556 (1983)), poly(2-hydroxyethyl methacrylate) (R. Langer et al., *J. Biomed. Mater. Res.*, **15**: 167-277 (1981), and R. Langer, *Chem. Tech.*, **12**: 98-105 (1982)), ethylene vinyl acetate (R. Langer et al., *Id.*) or poly-D-(-)-3-hydroxybutyric acid (EP 133,988).

Sustained release conjugate formulations also include liposomally entrapped conjugates. Such systems have the advantage that biologically active material can be introduced into tissues by phagocytosis, especially into tissues of the reticuloendothelial system. Liposomes containing conjugates are prepared by methods known *per se*, including: DE 3,218,121; Epstein et al., *Proc. Natl. Acad. Sci. U.S.A.*, **82**: 3688-3692 (1985); Hwang et al., *Proc. Natl. Acad. Sci. U.S.A.*, **77**: 4030-4034 (1980); G. Gregoriadis, *Liposome*

-16-

Technology, Vol. II. Incorporation of Drugs, Proteins, and Genetic Material, CRC Press, 1984; EP 52,322; EP 36,676; EP 88,046; EP 143,949; EP 142,641; Japanese Pat. Appln. 83-118008; U.S. Pat. Nos. 4,485,045; 4,774,085; and 4,544,545; and EP 102,324. In addition, the liposome may be antibody-coated for increase in uptake by the cells, as taught by Wang and Huang, Proc. Natl. Acad. Sci. U.S.A., **84**: 7851-7855 (1987). The liposomes obtained can be stored in the aqueous phase up to several weeks or months after addition of stabilizers, for example, lactose.

The size of the liposomes formed depends, for example, on the structure of the active ingredient and the lipid component, the mixing ratio of the lipid components, and the concentration of these components in the aqueous dispersion. Thus, for example, by increasing or reducing the concentration of the lipid component it is possible to produce aqueous phases having a high content of small or large liposomes. Ordinarily, the liposomes are of the small (about 200-800 Angstroms) unilamellar type in which the lipid content is greater than about 30 mol. percent cholesterol, the selected proportion being adjusted for the optimal therapy.

For parenteral administration, the liposome-containing aqueous dispersion is suitably mixed with customary thickeners, for example, hydroxypropylcellulose, suitable preservatives, and antioxidants, and can be used in the form of a lotion or gel for application to the skin or mucous membranes. For parenteral use, the aqueous dispersion of the enriched liposomes can be suspended in a suitable carrier liquid, for example, sterile, calcium free, isotonic sodium chloride or glucose solution, optionally buffered to pH 7.2-7.4.

It is estimated that the dose to be applied to a human of about 70 kg weight will range from about 200 mg to 1 g of liposomes containing about 0.1 to 500 mg of entrapped conjugate, respectively. However, the highest and lowest dose of the encapsulated material, the concentration of the phospholipids in the aqueous phase, as well as the proportions of the encapsulating phospholipids, can be varied according to results to be established experimentally in clinical trials, the "effective amount" being thereby tied to the therapeutic results so obtained.

The liposomal pharmaceutical administration system according to the present invention may consist of a kit of parts set comprising vials or bottles containing the phospholipids and conjugates.

If the antisense oligomer is to be used for cancer treatment, the conjugate therapy may be useful in conjunction with conventional chemotherapeutic agents, such as 5-fluorouracil. If the antisense oligomer is to be used for AIDS treatment, the conjugate therapy may be useful in conjunction with AZT, CD-4, and other experimental AIDS treatment agents.

If the conjugate herein is to be used for diagnostic purposes, it is labeled with a suitable label moiety, typically the oligonucleotide being labeled. Suitable labels include radioactive labels such as  $^{32}\text{P}$ ,  $^{125}\text{I}$ ,  $^{35}\text{S}$ , or the like, and non-radioactive labels such as, for example, biotin, thyroxine, enzymes such as hydrolases or peroxidases or phosphatases, and

-17-

various chemiluminescers such as luciferin, or fluorescent compounds such as fluorescein and its derivatives.

One diagnostic technique in which labeled conjugates are useful is in assays for nucleic acid having a predetermined nucleotide sequence in a sample relying on  
5 immobilization of the oligomeric probe on a solid support. Immobilizing the probe on a hydrophobic surface through the lipid moiety allows the oligonucleotide to be more accessible to hybridization than it is using, e.g., Southern blots or dot blots. Also, in the instant method, the filter need only be washed with a solution of the conjugate rather than heated to immobilize the nucleotides, as required for Southern blots.

10 The method herein involves immobilizing the conjugate, in labeled form, on a support, preferably a solid matrix such as an agarose gel or a filter membrane, and contacting the sample with the immobilized conjugate such that the nucleic acid will hybridize with the oligonucleotide portion of the conjugate if the nucleic acid is present in the sample. Conditions for hybridization are well known and include those described in  
15 Maniatis, Molecular Cloning: A Laboratory Manual (New York: Cold Spring Harbor Laboratory, 1982). Typically after such hybridization the support is washed to remove non-hybridizing materials and then the presence of labeled oligomers is determined.

In this context, the word "sample" refers to any liquid or biological sample that contains or potentially contains the nucleic acid to be assayed. Thus, this term includes  
20 fluids such as human or animal body fluids, e.g., blood, serum, urine, amniotic fluid, tissue extracts, cerebrospinal fluid, and the like.

The above diagnostic method is useful for detection of specific nucleic acid sequences associated with infectious diseases, genetic disorders, or cellular disorders such as cancer, e.g., oncogenes. Genetic diseases include specific deletions and/or mutations in  
25 genomic DNA from any organism, such as, e.g., sickle cell anemia, cystic fibrosis,  $\alpha$ -thalassemia,  $\beta$ -thalassemia, and the like. Infectious diseases can be diagnosed by the presence in clinic samples of specific DNA sequences characteristic of the causative microorganism. These include bacteria, such as Salmonella, Chlamydia, and Neisseria; viruses, such as the hepatitis viruses; and parasites, such as the Plasmodium responsible for  
30 malaria.

If a disease is characterized by the presence or absence of at least one specific restriction site in a specific nucleic acid sequence, such as sickle cell anemia and  $\beta$ -thalassemia, it may be detected by the use of restriction enzymes. Thus, in the above  
35 method, after the support is washed, the immobilized conjugate is treated with a restriction endonuclease so as to cleave a restriction site within the oligonucleotide portion of the conjugate, producing labeled and unlabeled oligomer fragments, as described in U.S. Pat. No. 4,725,537 issued February 16, 1988. The labeled oligomer fragments are then detected. Thus, DdeI [Geever et al., Proc. Natl. Acad. Sci., 78: 5081-5085 (1981)] or MstII [Orkin et al., N. Engl. J. Med., 307: 32-36 (1982)] may be employed if the disease is sickle cell  
40 anemia.

-18-

In another embodiment of diagnosis, a liposome incorporating the conjugate, prepared as described above, and a label (e.g., by encapsulation) is used in the detection of lupus erythematosus in humans. The method involves contacting serum from the human patient with the liposome (where the conjugate acts as the lupus-specific antigen) under conditions such that if lupus autoantibodies are present (as are contained in patients with active lupus), they will bind to the oligonucleotide on the liposome, thereby either stabilizing or destabilizing the liposome; and measuring for the presence or absence of label. If the binding to the oligonucleotide destabilizes the liposome, the label will be released and the amount of label released is measured. If the binding stabilizes the liposome, the label is absent and will not be detected. The assay generally involves one-minute incubation of the serum at room temperature and can be performed with standard microtiter plates, allowing the screening of large numbers of sera, in a short time. The conditions employed for incubation are suitably those used in the colorimetric test for lupus described by Janoff *et al.*, Clin. Chem., 29: 1587-1592 (1983) and EP 90,735 published Oct. 5, 1983. These references describe the liposome stabilization embodiment.

For the detection of lupus, preferably the oligonucleotide of the conjugate is double-stranded because it binds better to the antibodies. This may be accomplished by synthesizing a hairpin double-stranded nucleic acid, as is well known to those skilled in the art.

Another manner in which to use the conjugates herein is in the separation, isolation, or purification of a nucleic acid (e.g., oligonucleotides or polynucleotides) from a mixture of components. This procedure takes advantage of the altered physical properties of the oligonucleotide on the conjugate because of the presence of the lipid. The conjugate employed is one that has a nucleotide sequence on the oligonucleotide that is substantially complementary to a sequence contained within the nucleic acid being purified.

In one such use, the lipid of the conjugate is used to immobilize the oligonucleotide on a solid support containing a hydrophobic surface such as plastic, silanized glass, alkyl sepharose or other hydrophobic interaction chromatography support, or an RP-HPLC matrix (e.g., alkyl silica gels such as C8 Vydec). For this purpose, one need only dissolve the conjugate in a buffer such as a phosphate buffer and wash the resulting solution through the column. The lipid will adhere to the column under such aqueous conditions. Then the solid support is treated with the sample containing the complementary target nucleic acid, which hybridizes to the support under conditions that allow such hybridization as are well known to those skilled in the art. After hybridization the support is washed under different temperature or salt conditions than were employed to apply the mixture so as to denature the desired nucleic acid and isolate it. The conjugate can then be recovered from the column by washing it with acetonitrile or other organic solvent.

In another such use of this method to separate nucleic acids the lipid portion of the conjugate is used for transporting the complementary target nucleic acid through a water-non-miscible phase. Thus, the conjugate, having a nucleotide sequence capable of hybridizing to a sequence contained within the oligonucleotide to be separated, is provided

-19-

in a hydrophobic phase, such as octanol, in a suitable vessel for extraction and transfer from an organic phase to an aqueous phase, such as a U-tube. Likewise, the mixture containing the target nucleic acid is provided in a hydrophilic phase such as water in the vessel. The vessel also contains an aqueous phase without any nucleic acids, preferably  
5 pure water, into which the desired nucleic acid is transferred. The conjugate is partially soluble in both the aqueous and hydrophobic phases and its oligonucleotide portion would hybridize to the target strand, under suitable hybridizing conditions, and would then transport that strand through the hydrophobic phase into the pure hydrophilic phase. This procedure represents in effect a specific selective DNA extraction method.

10 The invention will be more fully understood by reference to the following examples. They should not, however, be construed as limiting the scope of the invention.

#### EXAMPLE I

##### Preparation and Purification of Conjugate

###### A. Preparation of Lipid Salt

15 1,2-di-O-hexadecyl-rac-glycerol (Sigma) in an amount of 342 mg (0.63 mmol) was evaporated from pyridine twice and then dissolved in 6 ml of pyridine and methylene chloride (1:1). After the flask was filled with argon, the solution, while being stirred at room temperature, was treated with van Boom's reagent (1.25 M solution in dioxane, 0.52 ml) (Marugg et al., Tetrahedron Lett., 27: 2661 (1986)). When the reaction was complete (as  
20 determined by thin layer chromatography using EM silica gel 60 F<sub>254</sub> pre-coated plates visualized with an acidic molybdenum stain where the starting material has an R<sub>f</sub> of 0.53 and the product has an R<sub>f</sub> of 0.24), the solution was diluted with methylene chloride (10 ml) and then shaken with several portions of aqueous triethylammonium bicarbonate (TEAB; 1 M). The organic layer was dried over anhydrous sodium sulfate, filtered, and  
25 evaporated in vacuo. Column chromatography performed with EM silica gel-60 (70-230 mesh) using SiO<sub>2</sub>, 5-20% methanol/methylene chloride with 1% added triethylamine followed by 1% acetic acid in place of triethylamine) followed by additional extractions of the purest fractions with 1 M TEAB afforded 1,2-di-O-hexadecyl-rac-glycero-3-H phosphonate as an off-white triethylammonium salt (76% yield).

30 <sup>1</sup>H NMR spectra were obtained using a Varian VXR-300S spectrometer and were recorded as ppm (d) using TMS as an internal standard. The results are <sup>1</sup>H NMR (d): 6.86 (d, J = 616 Hz, H-P); 3.90 (dd, J = 4.2 and 7.7 Hz, H<sub>2</sub>C-OP); 3.60-3.40 (m, 7 H, H<sub>2</sub>C-O-HC-H<sub>2</sub>C-O-H<sub>2</sub>C); 2.82 (qua, J = 7.2 Hz, 6H, (H<sub>2</sub>C)<sub>3</sub>-N); 1.54 (m, 4 H, 2(C-H<sub>2</sub>C-C-O)); 1.25 (bs, 26 H); 1.19 (t, J = 7.2 Hz, 9 H, (H<sub>3</sub>C-C)<sub>3</sub>-N); 0.87 (bt, J = 6.3 Hz, 6H, 2(H<sub>3</sub>C-C<sub>15</sub>)).

###### 35 B. Preparation of Conjugate

The synthesis of polymer-bound nucleotide H-phosphonates was performed on a Biosearch Model 4000 DNA synthesizer using a derivatized controlled-pore glass as the support in the method described in Froehler, B.C. and Matteucci, M.D., Tetrahedron Lett., 27: 469 (1986); Froehler, B.C. *et al.*, Nucl. Acids Res., 14: 5399 (1986); Froehler, B.C. and  
40 Matteucci, M.D., & Nucleotides, 6: 287 (1987); Garegg, P.J. *et al.*, Tetrahedron Lett., 27: 4051 (1986). Detritylation of the oligodeoxynucleotide afforded a free 5' hydroxyl group

-20-

suitable for conjugation with the lipid hydrogen phosphonate product of section A. This product was dissolved (25 mg/ml) in pyridine/acetonitrile (1:1) and conjugated with the polymer-bound oligodeoxynucleotide using pivaloyl chloride as the activating agent under the coupling conditions described in the above references. Doubling the final coupling time and/or repetition of the coupling cycle had little apparent effect on the yield or purity of the products. The compounds prepared were: EL-T<sub>10</sub>; EL-AGCTAGCT; EL-AGCTAGCTTTTGTAGCTAGCT; EL-CAGTGATGTGT; EL-ACACATCACTG, in which EL represents 1,2-di-O-hexadecyl-3-glycerolphosphate bound to the 5' hydroxyl group of the DNA.

After cleavage and deprotection using the methods in the above references, the products were analyzed using thin-layer chromatography, polyacrylamide gel electrophoresis, RP-HPLC, and enzymatic degradation.

#### C. Polyacrylamide gel electrophoresis

EL-DNA samples were dissolved in formamide-containing buffer, heated to 90°C for one minute, and loaded on 15% acrylamide, 7 M urea gels. After running at 50 mA (approximately 400 V) for 2.5 hours, gels were photographed under uv illumination. The lipid-containing EL-DNAs (visualized by UV shadowing) were less mobile than the corresponding lipid-free DNAs and appeared as broad streaks surrounding a major band. In each case, this major band was approximately as mobile as a normal oligomer twice as long as the DNA of the EL-DNA.

Polyacrylamide gel electrophoresis proved to be of little value, as the lipid-containing products appeared as indistinct smears of lower mobility than the corresponding non-lipid DNAs. Comparison of the relative amount of streaking EL-DNA to the normal pattern of DNA bands provided only a very rough measure of lipid incorporation.

#### D. Thin-layer chromatography

The samples were run on analytical TLC plates (SiO<sub>2</sub>, nPrOH:conc. aq. NH<sub>4</sub>OH:H<sub>2</sub>O 55:10:35) and visualized both by UV and staining (alcoholic p-anisaldehyde followed by heating).

Using the propanol/ammonia/water eluant, the different species were obtained as distinct, individual spots. The crude synthetic products typically showed two or three spots corresponding to EL-DNA, DNA, and sometimes a trace of unidentified material at lower R<sub>f</sub>. The major material (UV and stain) in all cases was the high R<sub>f</sub> (0.55-0.70) EL-DNA. The corresponding normal oligomers ran just above or just below the EL-DNAs. The relative locations of the compounds were constant, but their R<sub>f</sub> values were variable. In addition to the R<sub>f</sub> differences, the color of the spots (produced by reaction with the molybdenum stain) differed. EL-DNA appeared purple, while DNA was dark blue (as was the unidentified minor spot).

#### E. Reverse-phase HPLC

RP-HPLC was performed using a Waters Model 510 system and a 10 µm C8 Partisil-10, 4.6 x 250 mm analytical column. RP-HPLC samples were loaded in 100% "A" buffer (25 mM aqueous triethylammonium phosphate (TEAP), pH 7.0, 5% CH<sub>3</sub>CN). Increasing

-21-

amounts of "B" buffer (25 mM aqueous TEAP, pH 7.0, 75% CH<sub>3</sub>CN) were used to elute the samples from the column at a flow rate of 2.0 ml/min. Peaks were detected at 254 nm and were collected using an automated fraction collector.

5 Elution with the aqueous TEAP buffer with an acetonitrile gradient afforded good separation of EL-DNA from the small amount of contaminating normal DNA present in each reaction mixture. Under these conditions, DNA lacking the lipid eluted at 10-20% "B" depending on their length. In contrast to normal DNA, longer lipooligonucleotides were eluted before the shorter ones. Following are elution data for the five EL-DNAs (% "B"):  
10 EL-AGCTAGCTTTTAGCTAGCT (57%); EL-CAGTGATGTGT (62%); EL-ACACATCACTG (64%); EL-T<sub>10</sub> (65%); EL-AGCTAGCT (68%).

This technique gave the most detailed product information. Using TEAP buffer with an acetonitrile gradient, the presence or absence of lipid in the synthetic material was immediately obvious. In every case, there was a small amount of DNA (and the usual distribution of shorter oligomers) eluting at low acetonitrile percentages and a greater  
15 amount of late-eluting EL-DNA. A small analytical HPLC column generally produced a single, broad, tailing peak of lipid-DNA material. After the HPLC materials were isolated, further thin-layer chromatography experiments were run confirming that the late-eluting HPLC peaks and the high R<sub>f</sub>, purple-staining TLC spots were identical.

#### F. Enzymatic Digestion

20 Further characterization of one EL-DNA (EL-CAGTGATGTGT) was carried out by enzymatic digestion. Snake venom phosphodiesterase (Boehringer Mannheim GmbH) catalyzed nearly complete hydrolysis of the DNA backbone (36°C, 30 min., 50 mM Tris, pH 8, 10 mM MgCl<sub>2</sub>). Bovine spleen phosphodiesterase (Sigma), however, left the lipid-DNA intact (37°C, 1.5 hours, 50 mM Tris, pH 8, 10 mM MgCl<sub>2</sub>). These results are as expected  
25 for a DNA that bears a free hydroxyl group at the 3' terminus but is blocked at the 5' end. EL-CAGTGATGTGT was found to be incompetent as a substrate for both phospholipase C (Sigma, from *Bacillus cereus*) and phospholipase D (Sigma, from cabbage) under standard conditions (100 mM Tris, pH 8, 10 mM CaCl<sub>2</sub>, 36°C). RP-HPLC analysis of the digestion products showed only intact EL-DNA with each of the enzymes.

#### 30 G. Thermal Denaturation Profile

Samples containing normal DNA (CAGTGATGTGT/ACACATCACTG), EL-DNA (EL-CAGTGATGTGT/EL-ACACATCACTG), or a mixture of DNA/EL-DNA (EL-CAGTGATGTGT/ACACATCACTG and CAGTGATGTGT/EL-ACACATCACTG) in 1 ml of buffer (100 mM NaCl, 10 mM Na<sub>2</sub>HPO<sub>4</sub>, 1 mM EDTA, pH 7.2; each strand 2 μM)  
35 were placed in a masked 1-cm cuvette. All the samples were desalted (SEP-PAK C-18 minicolumn), HPLC-purified material. The insulated cell compartment was warmed from 10 to 80°C in 1°C increments with equilibration for 1 min. after attaining each temperature. UV absorbance at 260 nm was recorded automatically as a function of temperature.

The T<sub>m</sub>s for EL-CAGTGATGTGT/ACACATCACTG (38°C) and  
40 CAGTGATGTGT/EL-ACACATCACTG (33°C) were decreased relative to that of the normal duplex CAGTGATGTGT/ACACATCACTG (41°C). There have been a number

-22-

of reports of  $T_m$  decreases for DNA-ligand conjugates. See, for example, Kempe et al., Nucl. Acids Res., **13**: 45 (1985). The varying degree of  $T_m$  depression may simply reflect the differences in GC vs. AT content at the lipid-bearing end of each duplex. More interesting is that with lipid on both DNA strands (EL-CAGTGATGTGT/EL-  
5 ACACATCACTG;  $T_m$  50°C) the  $T_m$  is markedly higher than for lipid-free DNA. These results may indicate the formation of some higher-order lipid structures when lipid is present at each end of the duplex that are not accessible to duplexes bearing only a single lipid.

## EXAMPLE II

### 10 Transfection Assay of Conjugate

It is known that NIH/3T3 mouse fibroblasts can be transformed by ras oncogenes using calcium chloride transfection (see Graham and Van der Eb, Virology, **52**: 456-467 (1973) and Wigler et al., Proc. Natl. Acad. Sci. U.S.A., **76**: 1373-1376 (1979)). The transformed cells form foci of refringent cells that have lost contact inhibition and can  
15 grow in athymic mice.

In this prophetic example, the human ras oncogene (isolated from a plasmid, for example) is covalently coupled to a phospholipid by any of the techniques described above, including nitrogen linkage, or preferably is ligated to a lipid-oligonucleotide conjugate. In the latter case, a linker is constructed that is complementary to and spans the 3' end of the  
20 oligonucleotide of the conjugate to the 5' end of the human ras oncogene (which is contained in a linearized plasmid). This oligonucleotide portion of the conjugate, the linker, and the linearized plasmid are phosphorylated by standard kinase techniques and ligated together using T4 ligase.

Confluent monolayers of NIH/3T3 cells are washed twice with Dulbecco's Modified  
25 Eagle's medium (DMEM) without serum and incubated in this DMEM medium for four hours. Then, one of the lipid-ras conjugates is added to the medium and allowed to incubate for three hours at 37°C. Complete DMEM medium with 10% fetal calf serum is then added to the Petri dish, in an amount of 10:1 (v/v), and allowed to stand overnight at 37°C. The next day the cells are split in 1/20, 1/40, and 1/80 dilutions and left growing  
30 for 17 days, after which the foci are scored.

The ras oncogene can be internalized into the cell, brought to the nucleus, and expressed to yield the ras oncogene product, showing that the conjugates can transform NIH/3T3 cells in a stable fashion.

In summary, the present invention is directed to conjugates of oligonucleotides and  
35 lipids for transporting, targeting and internalizing the oligonucleotides to and within appropriate cells, generally by endocytosis. After internalization, the conjugate is cleaved, for example, by cellular lipases that recognize an appropriate cleavage site on the conjugate. Thus, the conjugate herein suitably acts as a cytospecific drug capable of specific delivery to a cell of an oligomer expressing a polypeptide having various types of activities, for  
40 example, therapeutic, prophylactic, antiviral, cytotoxic, etc. In addition, the oligomer may



-23-

affect the regulatory mechanism of the cell, complement a genetic defect, or serve as a marker.

-24-

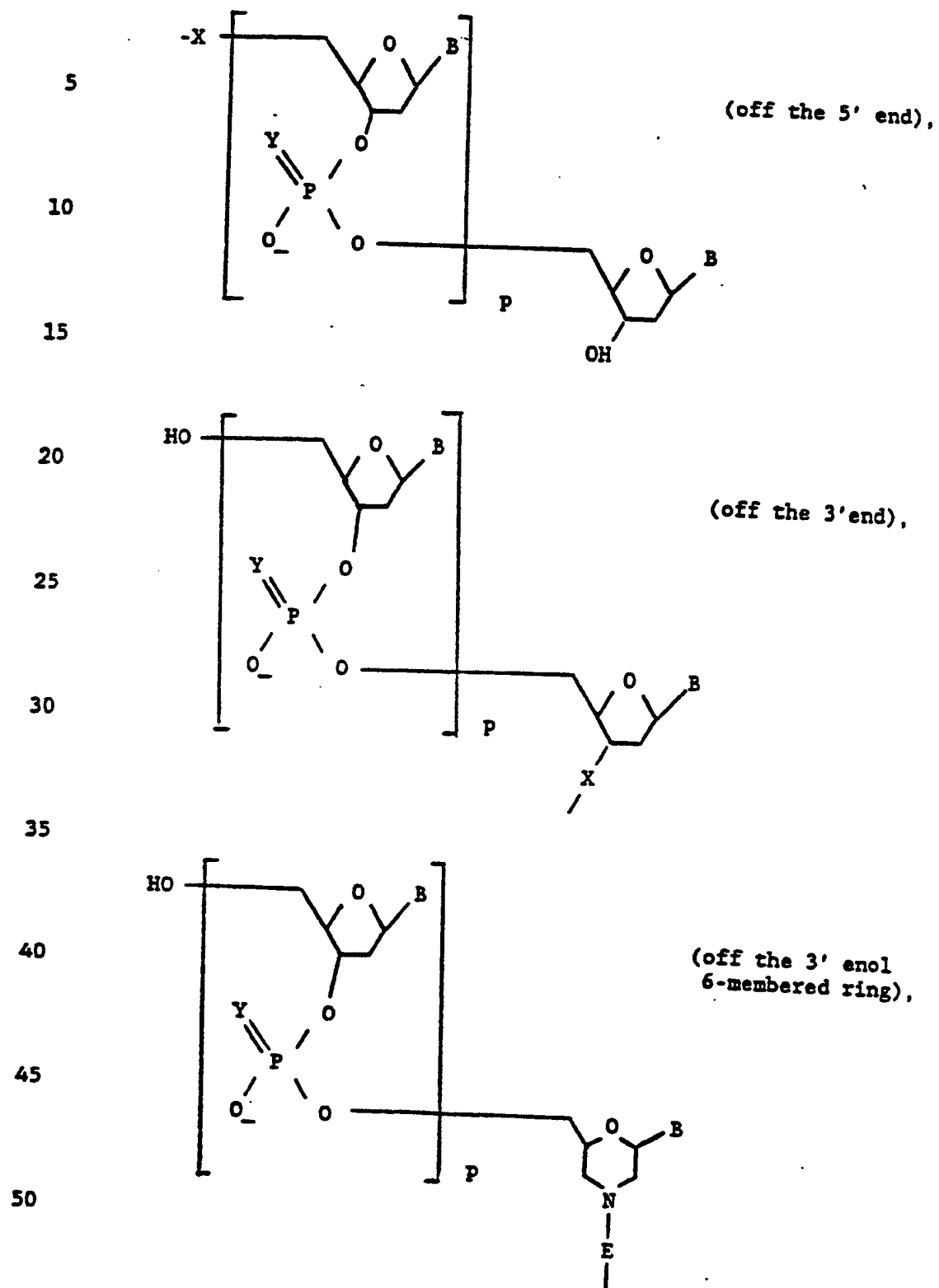
## WHAT IS CLAIMED IS:

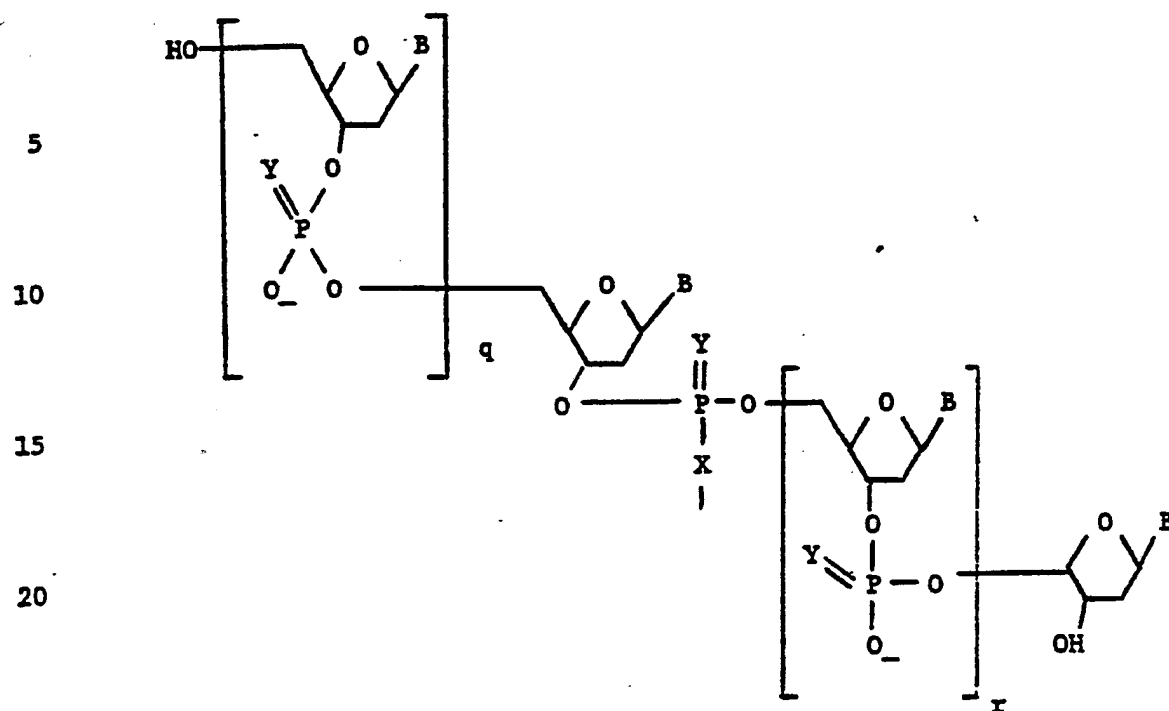
1. A covalent conjugate of a lipid and an oligonucleotide and pharmaceutically acceptable salts thereof.
2. The conjugate of claim 1 wherein the lipid is linked to the 5' end of the oligonucleotide.
3. The conjugate of claim 2 wherein the lipid is linked to the 5' hydroxyl group of the oligonucleotide.
4. The conjugate of claim 1 wherein the lipid has a cleavage site specifically recognized by an enzyme endogenous to a host cell to which the conjugate is targeted.
5. The conjugate of claim 4 wherein the cleavage site is susceptible to enzymatic hydrolysis.
6. The conjugate of claim 5 wherein the enzyme is located at a cellular or nuclear membrane.
7. The conjugate of claim 4 wherein the enzyme is in the cytoplasm of a cell.
8. The conjugate of claim 4 wherein the cleavage site is the oxygen atom derived from the 5' hydroxyl group of the oligonucleotide.
9. The conjugate of claim 3 wherein the lipid is a phospholipid and the enzyme is a lipase.
10. The conjugate of claim 9 wherein the lipase is a phospholipase.
11. The conjugate of claim 10 wherein the phospholipase is phospholipase C or D or both.
12. The conjugate of claim 11 wherein the cleavage site is the oxygen atom derived from the 5' hydroxyl group of the oligonucleotide.
13. The conjugate of claim 11 wherein the cleavage site is the oxygen atom on the glycerol side of the phosphoryl group of the phospholipid.
14. The conjugate of claim 1 having the formula:
 
$$L-([(X-Z)]_n-[X-P(=Y)O^{-1}_m])_y-A,$$
 wherein L is a steroid moiety,  $R_1$ , or  $R_2-X-CH(R_4)-CH(-X-R_3)-CH_2$ , where  $R_1$  is a  $C_1-C_{30}$  alkyl,  $C_2-C_{30}$  mono-, di-, or polyunsaturated alkyl,  $C_3-C_8$  cycloalkyl or  $C_4-C_8$  mono-, di-, or polyunsaturated cycloalkyl group,  $R_2$ ,  $R_3$ , and  $R_4$  are independently H,  $R_1$ , or  $\alpha$ -amino acyl, X is O, S, NH, C(=O), or C(=O)O, OC(=O), NHC(=O), or C(=O)NH, and Y is O or S;
 

Z is a  $C_2-C_{10}$  saturated or mono-, di-, or polyunsaturated alkylene moiety, n, m, and y are independently an integer from 0 to about 10; and

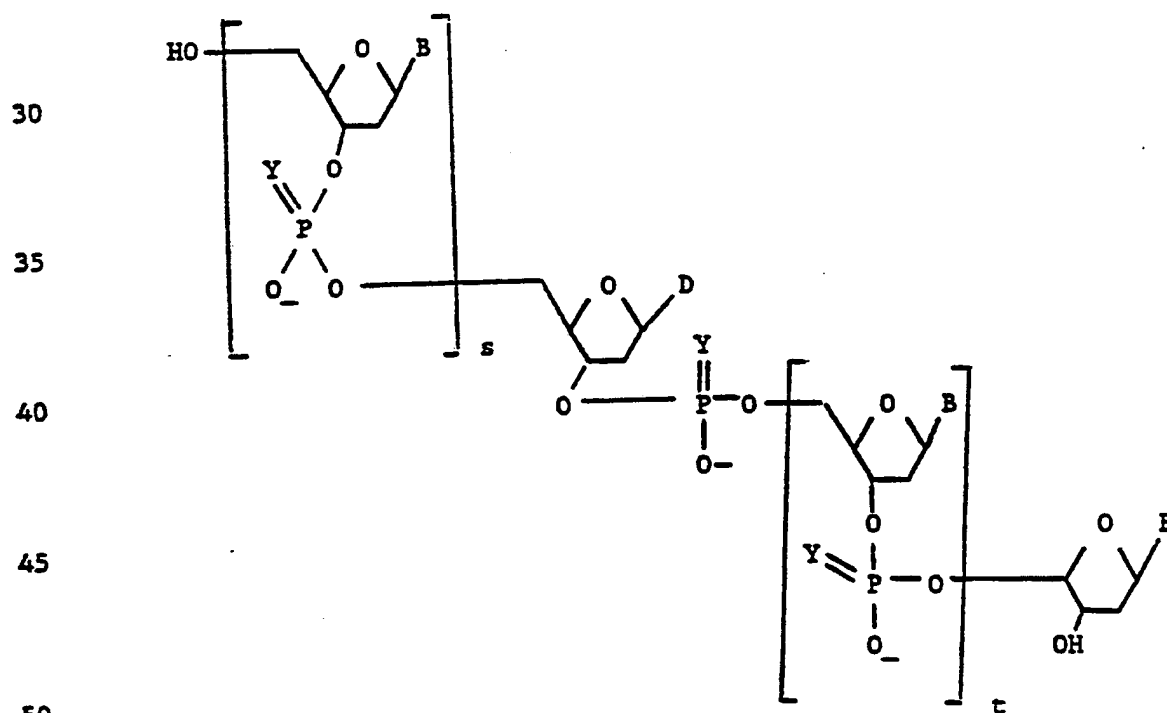
A is selected from the group consisting of

-25-



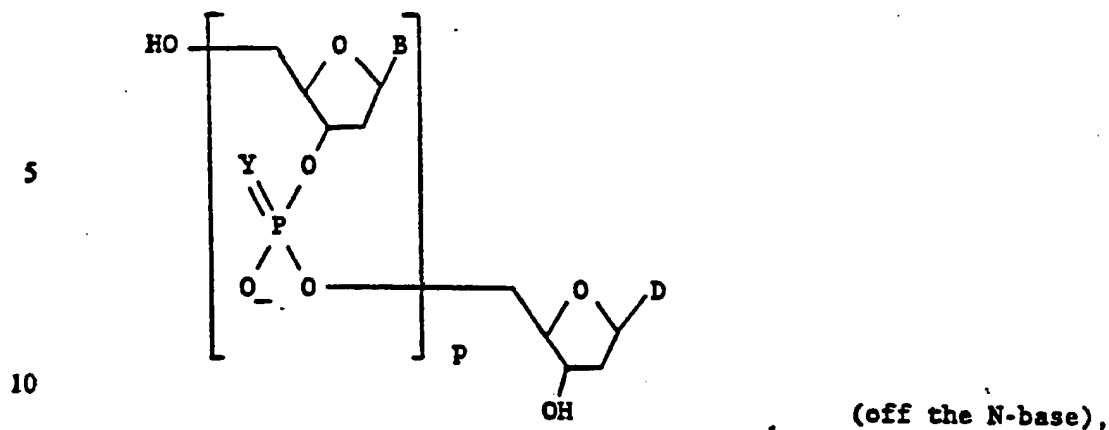


(off the internucleotidic phosphate),



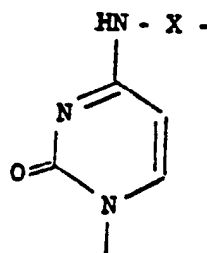
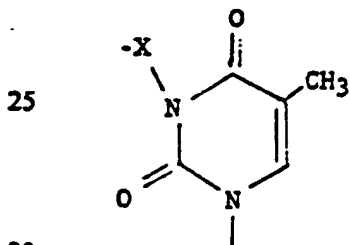
(off the N-base) , or

-27-

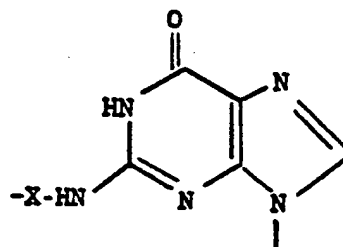
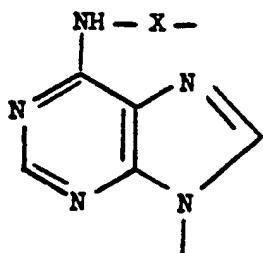


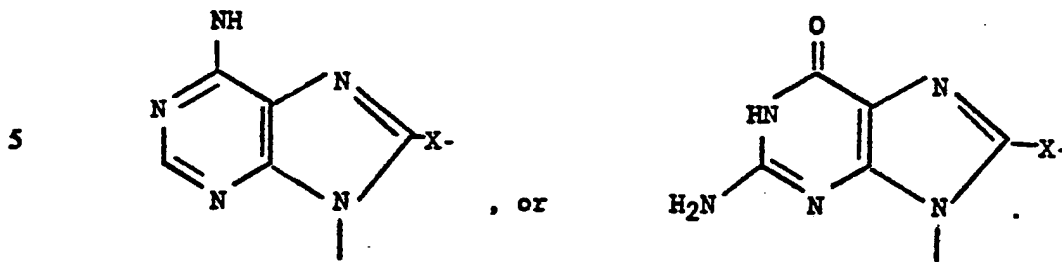
where B is a deprotected base, p is an integer from about 5 to 30, q and r are integers from about 1 to 28, provided that  $r + q$  is from about 4 to 29, s and t are integers from 0 to about 29, provided that  $s + t$  is from about 4 to 29, E is X or ZX, and D is selected from the group consisting of (where the bond from the ring nitrogens is attached to the sugar moiety of the oligonucleotide):

20



35





- 15 15. The conjugate of claim 14 wherein L is  $R_2-X-CH(R_4)-CH(-X-R_3)-CH_2$ .
16. The conjugate of claim 15 wherein  $R_2$  and  $R_3$  are independently H or  $C_{10} - C_{20}$  saturated or unsaturated alkyl groups,  $R_4$  is H or  $R_1$ , X is O, NH,  $NHC(=O)$ ,  $C(=O)$ ,  $OC(=O)$ , or  $C(=O)O$ , and Y is O.
17. The conjugate of claim 16 wherein Z is ethylene or carboxyethylene, n, m, and y are 0 to 2, and A is the oligonucleotide connected through its 5' end.
18. The conjugate of claim 17 where X is O, y is 1 or 2, m is 1 or 2, n is 0 or 1, and p is from about 10 to 25.
19. The conjugate of claim 15 wherein A is connected through the N6 of an adenine residue or the N4 of a cytosine residue.
20. The conjugate of claim 1 wherein the oligonucleotide comprises at least five bases.
- 25 21. The conjugate of claim 20 wherein the oligonucleotide has from about ten to about thirty bases.
22. The conjugate of claim 21 wherein the oligonucleotide has from about fourteen to about twenty-five bases.
23. The conjugate of claim 14 wherein at least one base is a substantially planar pyrimidinone base.
- 30 24. The conjugate of claim 1 wherein the oligonucleotide comprises a nucleotide sequence sufficiently complementary to a pathogenic nucleic acid or an oncogene to hybridize thereto.
25. The conjugate of claim 24 wherein the nucleic acid is viral nucleic acid.
- 35 26. The conjugate of claim 1 wherein the oligonucleotide has a nucleotide sequence representing, or capable of hybridizing to, a cleavage site specifically recognized by an enzyme endogenous to a host cell to which the conjugate is targeted.
27. The conjugate of claim 1 wherein the oligonucleotide comprises a nucleotide sequence representing, or capable of hybridizing to, an mRNA splice site.
- 40 28. The conjugate of claim 1 that is labeled.
29. The conjugate of claim 1 that is immobilized on a solid support.
30. The conjugate of claim 29 wherein the solid support has a hydrophobic surface.
31. A method comprising transfecting into a host cell the conjugate of claim 1.
- 45 32. A host cell transfected with the conjugate of claim 1.

-29-

33. A composition comprising a pharmaceutically acceptable carrier and the conjugate of claim 1.
34. The composition of claim 33 wherein the carrier is a liposome.
35. The composition of claim 33 that is isotonic.
- 5 36. The composition of claim 33 that is sterile.
37. A method comprising administering to a mammal suffering from a pathogenic condition an effective amount of the composition of claim 33.
38. The method of claim 37 wherein the pathogenic condition is a viral infection.
39. The method of claim 37 wherein the carrier in the composition is a liposome.
- 10 40. A method for the assay of a nucleic acid having a predetermined nucleotide sequence in a sample comprising:
- (a) providing the conjugate of claim 28 that has a nucleotide sequence capable of hybridizing to the predetermined sequence;
  - (b) immobilizing the labeled conjugate on a support;
  - 15 (c) contacting the sample with the immobilized conjugate under conditions that would cause hybridization of the nucleic acid with the oligonucleotide portion of the conjugate if the nucleic acid is present in the sample; and
  - (d) detecting the presence of labeled oligomers.
- 20 41. The method of claim 40 which further comprises, after step (c), the step of contacting the immobilized conjugate under conditions of digestion with a restriction endonuclease that is capable of cleaving the oligonucleotide portion of the conjugate so as to produce labeled and unlabeled oligomer fragments; and wherein in step (d) the oligomers are oligomer fragments.
- 25 42. A method for separating an oligonucleotide from a mixture, which method comprises:
- (a) providing the conjugate of claim 1 having a nucleotide sequence capable of hybridizing to a sequence contained within the oligonucleotide to be separated;
  - (b) immobilizing the conjugate on a support;
  - (c) contacting the mixture with the immobilized conjugate, under conditions
  - 30 causing hybridization of the oligonucleotide of the mixture with the oligonucleotide portion of the conjugate; and
  - (d) separating the hybridized oligonucleotides.
43. A method for separating an oligonucleotide from a mixture, which method comprises:
- 35 (a) providing the conjugate having a nucleotide sequence capable of hybridizing to a sequence contained within the oligonucleotide to be separated in a hydrophobic phase;
  - (b) providing the mixture in a hydrophilic phase;
  - (c) contacting the phase containing the mixture with the phase containing the
  - 40 conjugate, under conditions causing hybridization of the oligonucleotide of the mixture with the oligonucleotide portion of the conjugate; and

-30-

(d) extracting from the phase containing the mixture the hybridized oligonucleotides and transporting them to a separate hydrophilic phase.

44. A liposome comprising the conjugate of claim 1 and a label moiety.

5 45. A method for detecting lupus erythematosus in a human comprising contacting a serum sample from the human with the liposome of claim 44, said contacting being such that the antibodies bind to the oligonucleotide on the liposome, so as to alter the stability of the liposome; and measuring for the presence or absence of the label.

10 46. The method of claim 45 wherein the antibodies bind to the oligonucleotide on the liposome so as to release the label and wherein the amount of the label released is measured.





## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

|  |           |   |
|--|-----------|---|
| <b>(51) International Patent Classification 5 :</b><br><b>A61K 47/48, 31/70</b>  | <b>A3</b> | <b>(11) International Publication Number:</b> <b>WO 90/10448</b><br><b>(43) International Publication Date:</b> 20 September 1990 (20.09.90)  |
| <b>(21) International Application Number:</b> PCT/US90/01002<br><b>(22) International Filing Date:</b> 23 February 1990 (23.02.90)<br><b>(30) Priority data:</b><br>320,202 7 March 1989 (07.03.89) US<br><b>(71) Applicant:</b> GENENTECH, INC. [US/US]; 460 Point San Bruno Boulevard, South San Francisco, CA 94080 (US).<br><b>(72) Inventors:</b> BISCHOFBERGER, Norbert, W. ; 1116 Cedar Street, San Carlos, CA 94070 (US). SHEA, Regan, G. ; 365 Laurie Meadows Drive No. 219, San Mateo, CA 94403 (US).<br><b>(74) Agents:</b> WINTER, Daryl, B. et al.; Genentech, Inc., 460 Point San Bruno Boulevard, South San Francisco, CA 94080 (US). |           | <b>(81) Designated States:</b> AT (European patent), BE (European patent), CA, CH (European patent), DE (European patent), DK (European patent), ES (European patent), FR (European patent), GB (European patent), IT (European patent), JP, LU (European patent), NL (European patent), SE (European patent).<br><br><b>Published</b><br><i>With international search report.</i><br><i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i><br><br><b>(88) Date of publication of the international search report:</b><br>29 November 1990 (29.11.90) |
| <b>(54) Title:</b> COVALENT CONJUGATES OF LIPID AND OLIGONUCLEOTIDE<br><br><b>(57) Abstract</b><br><br>A conjugate is provided wherein a lipid is covalently coupled to an oligonucleotide having a nucleotide sequence that is either of interest or substantially complementary to a nucleotide sequence of interest. The conjugate is useful for nucleic acid purification, diagnostic methods, and therapeutic applications. The therapeutic aspect includes conjugates wherein the oligonucleotide contains an antisense nucleotide sequence to a sequence encoding a polypeptide responsible for a pathogenic disorder.                        |           |   |

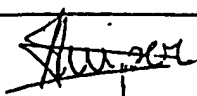
**FOR THE PURPOSES OF INFORMATION ONLY**

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

|    |                              |    |  |    |                          |
|----|------------------------------|----|--|----|--------------------------|
| AT | Austria                      | ES | Spain                                    | MC | Monaco                   |
| AU | Australia                    | FI | Finland                                  | MG | Madagascar               |
| BB | Barbados                     | FR | France                                   | ML | Mali                     |
| BE | Belgium                      | GA | Gabon                                    | MR | Mauritania               |
| BF | Burkina Faso                 | GB | United Kingdom                           | MW | Malawi                   |
| BG | Bulgaria                     | GR | Greece                                   | NL | Netherlands              |
| BJ | Benin                        | HU | Hungary                                  | NO | Norway                   |
| BR | Brazil                       | IT | Italy                                    | RO | Romania                  |
| CA | Canada                       | JP | Japan                                    | SD | Sudan                    |
| CF | Central African Republic     | KP | Democratic People's Republic<br>of Korea | SE | Sweden                   |
| CG | Congo                        | KR | Republic of Korea                        | SN | Senegal                  |
| CH | Switzerland                  | LI | Liechtenstein                            | SU | Soviet Union             |
| CM | Cameroon                     | LK | Sri Lanka                                | TD | Chad                     |
| DE | Germany, Federal Republic of | LU | Luxembourg                               | TG | Togo                     |
| DK | Denmark                      |    |  | US | United States of America |

# INTERNATIONAL SEARCH REPORT

International Application No PCT/US 90/01002

|  |  |  |
|--|--|--|
| <b>I. CLASSIFICATION OF SUBJECT MATTER</b> (if several classification symbols apply, indicate all) <sup>6</sup>  |  |  |
| According to International Patent Classification (IPC) or to both National Classification and IPC  |  |  |
| IPC <sup>5</sup> : A 61 K 47/48, A 61 K 31/70  |  |  |
| <b>II. FIELDS SEARCHED</b>   |  |  |
| Minimum Documentation Searched <sup>7</sup>  |  |  |
| Classification System  | Classification Symbols   |  |
| IPC <sup>5</sup>   | A 61 K, C 07 H   |  |
| Documentation Searched other than Minimum Documentation<br>to the extent that such Documents are included in the Fields Searched <sup>8</sup>  |  |  |
|  |  |  |
| <b>III. DOCUMENTS CONSIDERED TO BE RELEVANT <sup>9</sup></b>   |  |  |
| Category <sup>9</sup>  | Citation of Document, <sup>11</sup> with indication, where appropriate, of the relevant passages <sup>12</sup>   | Relevant to Claim No. <sup>13</sup>  |
| X  | Chemical Reviews, vol. 90, no. 4, June 1990, American Chemical Society, E. Uhlmann et al.: "Antisense oligo-nucleotides: a new therapeutic principle", pages 556-562<br>see page 560, left-hand column, paragraph d)<br>& Human Retroviruses, Cancer and Aids, Approaches to Prevention and Therapy, (Goodchild et al.) 1988, New York, page 423,... | 1-36, 40-42, 44-46   |
| X  | STN File Server (Karlsruhe) & File Medline, H. Yanagawa et al.: "Analysis of super-helical structures of nucleic acid-lipid conjugates by image processing", see accession number 89144950, & Nucleic Acids Symp. Ser., (1988), (19), 189-92   | 1-36, 40-42, 44-46   |
|  | --   |  |
|  | ./.  |  |
| <div style="display: flex; justify-content: space-between;"> <div style="width: 45%;"> <p><sup>10</sup> Special categories of cited documents:</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> </div> <div style="width: 45%;"> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.</p> <p>"Z" document member of the same patent family</p> </div> </div> |  |  |
| <b>IV. CERTIFICATION</b>   |  |  |
| Date of the Actual Completion of the International Search<br>6th August 1990   |  | Date of Mailing of this International Search Report<br>29 OCT 1990   |
| International Searching Authority<br><br>EUROPEAN PATENT OFFICE  |  | Signature of Authorized Officer<br><br>Mme N. KUIPER  |

| III. DOCUMENTS CONSIDERED TO BE RELEVANT (CONTINUED FROM THE SECOND SHEET) |   |                       |
|--|---|-----------------------|
| Category *   | Citation of Document, " with indication, where appropriate, of the relevant passages  | Relevant to Claim No. |
| X  | Nucleic Acids Research, vol. 15, no. 14, 1987, IRL Press Ltd, (Oxford, GB),<br>C.J. Marcus-Sekura et al.: "Comparative inhibition of chloramphenicol acetyltransferase gene expression by antisense oligonucleotide analogues having alkyl phosphotriester, methylphosphonate and phosphorothioate linkages", pages 5749-5763<br>see figure 1; isopropyl and ethylphosphotriester<br>cited in the application<br>-- | 1-36,40-42,<br>44-46  |
| X  | Biotechniques, vol. 6, no. 10, 1988,<br>A.R. van der Krol et al.: "Modulation of eukaryotic gene expression by complementary RNA or DNA sequences", pages 958-974<br>see table 2<br>cited in the application<br>--  | 1-36,40-42,<br>44-46  |
| X  | STN File Server (Karlsruhe) & File CA, Chemical Abstracts, vol. 109, 1988, (Columbus, Ohio; US),<br>E.E. Bichenkov et al.: "Interaction of cholesterol-modified polynucleotide with phosphatidylcholine liposomes", see abstract 225156z,<br>& Biol. Membr., 5(7), 735-42<br>--   | 1-36,40-42,<br>44-46  |
| X  | DE, A, 3637243 (TOAGOSEI CHEM. IND.)<br>7 May 1987<br>see claims<br>--  | 1-36,40-42,<br>44-46  |
| P,X  | Proc. Natl. Acad. Sci. USA, vol. 86, September 1989,<br>R.L. Letsinger et al.: "Cholesteryl-conjugated oligonucleotides: Synthesis, properties, and activity as inhibitors of replication of human immunodeficiency virus in cell culture", pages 6553-6556<br>see the whole article<br>-----   | 1-36,40-42,<br>44-46  |

## FURTHER INFORMATION CONTINUED FROM THE SECOND SHEET

V. ☒ OBSERVATIONS WHERE CERTAIN CLAIMS WERE FOUND UNSEARCHABLE <sup>1</sup>

This international search report has not been established in respect of certain claims under Article 17(2) (a) for the following reasons:

1. ☒ Claim numbers ..... \* because they relate to subject matter not required to be searched by this Authority, namely:

\* 37-39

See PCT Rule 39.1(iv): methods for treatment of the human or animal body by surgery or therapy, as well as diagnostic methods

2. ☐ Claim numbers ....., because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

3. ☐ Claim numbers ....., because they are dependent claims and are not drafted in accordance with the second and third sentences of PCT Rule 6.4(a).

VI. ☒ OBSERVATIONS WHERE UNITY OF INVENTION IS LACKING <sup>2</sup>

This international Searching Authority found multiple inventions in this international application as follows:

1. claims 1-36, 40-42 and 44-46 (all in part)
2. claims 1, 4-8, 10-16, 19-36, 40-42 and 44-46 (all in part)
3. claims 1-36, 40-42 and 44-46 (all in part)
4. claims 1, 4-8, 10-16, 19-36, 40-42 and 44-46 (all in part)
5. claim 43

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims of the international application.
2. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims of the international application for which fees were paid, specifically claims:

3. ☒ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claim numbers:

1-36, 40-42 and 44-46 (all in part)

4. ☐ As all searchable claims could be searched without effort justifying an additional fee, the International Searching Authority did not invite payment of any additional fee.

## Remark on Protest

- ☐ The additional search fees were accompanied by applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

US 9001002  
SA 35106

**This annex lists the patent family members relating to the patent documents cited in the above-mentioned international search report. The members are as contained in the European Patent Office EDP file on 23/10/90. The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.**

| Patent document<br>cited in search report | Publication<br>date | Patent family<br>member(s) | Publication<br>date |
|---|---------------------|----------------------------|---------------------|
| DE-A- 3637243                             | 07-05-87            | JP-A- 63030497             | 09-02-88            |
|   |                     | JP-A- 63039894             | 20-02-88            |
|   |                     | US-A- 4808708              | 28-02-89            |
|   |                     | JP-A- 62201895             | 05-09-87            |
| <hr/>                                     |                     |                            |                     |

**This Page is Inserted by IFW Indexing and Scanning  
Operations and is not part of the Official Record**

## **BEST AVAILABLE IMAGES**

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images include but are not limited to the items checked:

- ☐ BLACK BORDERS
- ☒ IMAGE CUT OFF AT TOP, BOTTOM OR SIDES
- ☐ FADED TEXT OR DRAWING
- ☐ BLURRED OR ILLEGIBLE TEXT OR DRAWING
- ☐ SKEWED/SLANTED IMAGES
- ☐ COLOR OR BLACK AND WHITE PHOTOGRAPHS
- ☐ GRAY SCALE DOCUMENTS
- ☒ LINES OR MARKS ON ORIGINAL DOCUMENT
- ☐ REFERENCE(S) OR EXHIBIT(S) SUBMITTED ARE POOR QUALITY
- ☐ OTHER: \_\_\_\_\_

**IMAGES ARE BEST AVAILABLE COPY.**

**As rescanning these documents will not correct the image problems checked, please do not report these problems to the IFW Image Problem Mailbox.**